

SHANKS & HERBERT

TransPotomac Plaza
1033 N. Fairfax St.
Suite 306
Alexandria, VA 22314
(703) 683-3600

1c803 U.S. PTO



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Inventor: David M. Margolis, et al.For: An HIV Transcription Repressor And Compositions and Methods Based Thereon

Enclosed are:

- ☒ 53 pages of specification; 3 pages of claims (claims 1-23); a one (1) page abstract; and 18 sheets of drawings (Figs. 1a-10b).
- ☐ An assignment of the invention to _____
- ☐ A certified copy of a _____ application
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Respectfully submitted,

SHANKS & HERBERT

Chalin A. Smith
Reg. No. 41,569

UNITED STATES PATENT APPLICATION

For:

**AN HIV TRANSCRIPTION REPRESSOR COMPLEX
AND COMPOSITIONS AND METHODS BASED THEREON**

Inventors:

David M. Margolis

Fabio Romerio

Jason J. Coull

University of Maryland Biotechnology Institute
Institute of Human Virology
University of Maryland
725 W. Lombard Street
Baltimore, MD 21201

**AN HIV TRANSCRIPTION REPRESSOR COMPLEX AND COMPOSITIONS
AND METHODS BASED THEREON**

TECHNICAL FIELD OF INVENTION

The present invention relates to our discovery of the molecular mechanism by which a protein complex inhibits HIV-1 transcription. More particularly, the invention relates to a transcription repressor complex or "TRC" containing YY1, LSF, and HDAC1. The components of the TRC cooperate to uniquely bind to at least a portion of the long term repeat (LTR) sequence of HIV-1, preferably the repressor complex sequence (RCS) and inhibit transcription. The invention further relates to pharmaceutical compositions that manipulate this mechanism and methods of use thereof to treat both active and latent HIV infection.

CROSS REFERENCE TO RELATED APPLICATION

This application is a continuation-in-part of U.S. Patent Application Serial. No. 09/355,010 filed July 22, 1999, which in turn claims the benefit of PCT Application Serial No. PCT/US98/00574 (published as International Publication No. WO 98/33067), filed January 13, 1998, which in turn claims the benefit of provisional application Serial No. 60/036,242, filed January 23, 1997, now abandoned, all of which are incorporated by reference herein in their entirety.

ACKNOWLEDGEMENT

This work was supported in part by a grant from the Federal Government, National Institutes of Health grant number AI 41366 and AI 45297. The Government therefore has certain rights in the invention.

BACKGROUND OF THE INVENTION

Human immunodeficiency virus is a retrovirus whose major cell target is CD4⁺ T-lymphocytes. HIV-1 infection is mediated through the binding of the virus to the CD4 glycoprotein and other co-receptors. The HIV-1 envelope glycoproteins gp41 and gp120 direct this binding. The gp120 is non-covalently attached to gp41, which is anchored in the viral lipid bilayer. HIV-1 entry is mediated by the high-affinity

binding of gp120 to the amino-terminal domain of the CD4 glycoprotein, causing conformational changes in gp120 (McDougal *et al.*, 1986, *Science* 231:382-385; Helseth *et al.*, 1990, *J. Virol.* 64:2416-2420; Wain-Hobson, 1996, *Nature* 384:117-118) and subsequent binding of gp120 to co-receptors, such as CXCR4 and CC-CKR5 (Wu *et al.*, 1996, *Nature* 384:179; Trkola *et al.*, 1996, *Nature* 384:184; Wain-Hobson, 1996, *Nature* 384:117-118).

Studies have shown that the LTR, the promoter of HIV-1, responds to numerous cellular factors (Garcia *et al.*, 1987, *EMBO J.* 6:3761-3770; Giacca *et al.*, 1992, *Virol.* 186:133-147; Harrich *et al.*, 1989, *J. Virol.* 63:2858-2891; Jones, 1989, *New Biol.* 1:127-135; Jones *et al.*, 1986, *Science* 232:755-759; Nabel and Baltimore, 1988, *Nature* 326:711-713; and Wu *et al.*, 1988, *J. Virol.* 62:218-225). Such factors may augment or repress the production of HIV-1 virions by infected cells.

Two cellular factors, YY1 and LSF that cooperate uniquely in recognition of the region -10 to +27 of the HIV-1 LTR (referred to as the "RCS", for repressor complex sequence) have previously been identified. These have been shown to specifically and synergistically repress HIV LTR expression and viral production (41, 49). Antibodies to either YY1 or LSF inhibit RCS formation, and mutations within the LTR that eliminate LSF binding and RCS formation ablate repression mediated by YY1 and/or LSF (49).

Yin Yang 1 or "YY1" is a widely distributed 68 kDa multifunctional transcriptional regulator with homology to the GLI-Krüppel family of proteins. YY1 has the ability both to activate and repress gene expression (16, 32, 52, 56). It contains a plurality of "zinc fingers" as well as two N-terminal transactivation domains, while the C-terminal domain is required for direct DNA binding and for repression of some promoters (2, 4, 17). This broad spectrum of activity has been attributed to bending of DNA, interactions with other factors, or post-transcriptional modification of YY1 (52). However, activity depends on the promoter context and specific protein-protein interactions that YY1 establishes with other regulatory proteins (23, 32-34, 49, 50, 53, 70-72, 76), and with general transcription factors (5, 60).

"YY1" directly interacts with many viral and cellular nuclear factors (Shi *et al.*, 1991, *Cell* 67:377-388; Lee *et al.*, 1993, *Proc. Natl. Acad. Sci. USA* 90:6145-6149; Chiang *et al.*, 1995, *Science* 267:531-536; Zhou *et al.*, 1995, *J. Virol.* 69:4323-

4330). YY1 has been shown to regulate both viral and lymphocyte promoters (Bauknecht *et al.*, 1992, EMBO J. 11:4607-4617; Flanagan *et al.*, 1992, Mol. Cell Biol. 12:38-44; Park and Atchison, 1991, Proc. Natl. Acad. Sci. USA 88:9804-9808; Seto *et al.*, 1991, Nature 354:241-245; and Shi *et al.*, 1991, Cell 67:377-388).

5 YY1 (also known as δ , NF-E1, UCRBP or CF1) has been shown to cooperate uniquely in recognition of the region of the HIV-1 LTR referred to as the repressor complex sequence or RCS (Malim *et al.*, 1989, J. Virol. 63:3213-9; Shi *et al.*, 1997, Biochim Biophys Acta. 1332:F49-66; Shrivastava, *et al.*, 1994, Nucleic Acids Res. 22:5151-5; Wu, *et al.*, 1988, EMBO J. 7:2117-30). YY1 has previously been shown
10 to repress HIV-1 transcription and virion production (Margolis, *et al.*, 1994, J. Virol. 68:905-910). Moreover YY1 had been found to inhibit HIV-1 LTR transcription *in vivo* (Margolis *et al.*, 1994, J. Virol. 68:905-910).

LSF is a lymphoid transcription factor that has been shown to repress LTR transcription in *in vitro*, but not in *in vivo* assays (Kato *et al.*, 1991, Science 251:1476-
15 1479; Yoon *et al.*, 1994, Mol. Cell. Biol. 14:1776-1785). LSF is the predominantly expressed member of a family of proteins (also termed LBF-1, CP-2, LBP-1a, b, c and d) that are produced from the differential splicing from two related genes (55, 73). All bind DNA except for LSF-ID (LBP1-d), which lacks a central encoding exon. LSF (LBF-1, CP-2) recognizes the same LTR sequence as YY1 (Huan *et al.*, 1990, Genes
20 Dev. 4:287-298; Lim *et al.*, 1992, Mol. Cell. Bio. 12:828-835; Garcia *et al.*, 1987, EMBO J. 6:3761-3770; Kato *et al.*, 1991, Science 251:1476-1479; Yoon *et al.*, 1994, Mol. Cell. Biol. 14:1776-1785). LSF binding to the LTR is associated with direct repression of transcription *in vitro* (18, 29, 44). However, transient expression of LSF alone had no observable effect on expression from the HIV LTR (49, 73, 75).

25 As we reported in parent application, Serial No. 09/355,010 (International Publication No. WO 98/33067), the entire contents of which are incorporated by reference herein, YY1 and LSF form a complex (the TRC) that binds the HIV LTR, particularly the RCS site, and blocks transcription. As discussed in detail herein, we have discovered that the complex further contains HDAC1. A molecular mechanism
30 of TRC-associated repression of HIV-1 which is also described herein. The invention is based on the discovery that (1) HDAC1 copurifies with the LTR-binding YY1-LSF repressor complex; (2) the domain of YY1 that interacts with HDAC1 is required to

repress the HIV-1 promoter; (3) the expression of HDAC1 augments repression of the LTR by YY1, and (4) the deacetylase inhibitor trichostatin-A blocks repression mediated by YY1. This novel link between HDAC1 recruitment and inhibition of HIV-1 expression by YY1 and LSF, in the natural context of a viral promoter integrated into chromosomal DNA, supports novel HIV therapies described herein and has significant implications for the long-term treatment of AIDS.

A subpopulation of stably infected CD4⁺ T lymphocytes containing integrated proviral DNA capable of producing virus upon stimulation has been identified in HIV⁺ individuals (6, 7, 8, 15, 68). As antiretroviral therapy now allows significant inhibition of active HIV-1 replication, an understanding of factors that establish or maintain the integrated proviral state takes on new relevance. Potent repression of LTR transcription could allow an activated, infected cell to return to the resting state and establish stable nonproductive infection. This may occur via changes in local chromatin architecture surrounding the HIV promoter. While activation of the HIV LTR has been shown to be associated with changes in chromatin structure (13, 46, 51, 61-63), factors that result in durable repression of LTR expression are less well known.

In view of the lack of methods of down-regulating transcription of the HIV-1 LTR, it is clear that there exists in the art a need for effective therapies that regulate transcriptional repression of the HIV-1 LTR. The present invention solves this problem by providing methods for the improvement of HIV-1 LTR transcriptional repression. It also solves this problem by providing methods for improved antagonism of LTR transcriptional repression, thereby leading to conditions in which HIV cannot establish a virologically latent intracellular infection, and that will allow for clearance of HIV infection when used in combination with other potent anti-viral agents. The regulation of proviral expression within this reservoir of infected CD4⁺ cells may take on new relevance as potent combination antiretroviral therapies allow the depletion of HIV from productively infected cell populations.

SUMMARY OF THE INVENTION

The present invention relates to our discovery of the molecular mechanism by which the YY1/LSF repressor complex inhibits HIV-1 transcription. Our discovery of

a novel link between HDAC1 recruitment and inhibition of HIV-1 expression by YY1 and LSF, in the natural context of a viral promoter integrated into chromosomal DNA, supports novel HIV therapies described herein and has significant implications for the long-term treatment of AIDS.

5 An object of the present invention is to provide a method of repressing HIV transcription, thereby treating or preventing HIV infection in a human subject in need of such treatment.

10 In one preferred embodiment, this method comprises administering to the subject an effective amount of a preparation of a transcription repressor complex comprising YY1 (or a derivative or analog thereof), LSF (or a derivative or analog thereof), and HDAC1 (or a derivative or analog thereof). Preferred derivatives and analogs are discussed in detail herein.

15 In another preferred embodiment, this method comprises administering to the subject an effective amount of a preparation comprising an agent that enhances the binding of a YY1 (or a derivative or analog thereof) to HDAC1 (or a derivative or analog thereof).

20 In another preferred embodiment, this method comprises administering to the subject an effective amount of an agent that enhances recruitment of HDAC1 (or a derivative or analog thereof) by YY1 (or a derivative or analog thereof) to the LTR RCS site.

 In another preferred embodiment, this method comprises administering to the subject an effective amount of a preparation comprising an agent that enhances the activity of HDAC1, or a derivative or analog thereof.

25 In another preferred embodiment, this method comprises administering to the subject an effective amount of an agent that enhances the expression of HDAC1, or a derivative or analog thereof.

 In another preferred embodiment, this method comprises administering to the subject an effective amount of an agent that up-regulates the expression of HDAC1, or a derivative or analog thereof.

30 In another preferred embodiment, this method comprises administering to the subject an effective amount of a nucleic acid or combination of nucleic acids comprising: one or more nucleotide sequences encoding YY1 (or a derivative or

analog thereof); one or more nucleotide sequences encoding LSF (or a derivative or analog thereof); and one or more nucleotide sequences encoding HDAC1 (or a derivative or analog thereof).

5 A further object of the present invention is to provide a method of treating quiescent reservoirs of HIV infection in a human subject in need of such treatment comprising the steps of:

- 10 (a) administering to the subject an amount of an agent that down-regulates the expression of HDAC1, or a derivative or analog thereof, the amount effective to down-regulate TRC-associated repression of HIV transcription;
- (b) allowing latent, quiescent reservoirs of HIV to become actively transcribing; and
- (c) treating the subject with an effective amount of an antiretroviral agent.

15 In an alternate embodiment, this method comprises the administration of an agent that inhibits the expression of HDAC1 or a derivative or analog thereof, the amount being effective to inhibit TRC-associated repression of HIV transcription.

20 Another object of the present invention is to provide a composition comprising the TRC complex of YY1, LSF and HDAC1 and a method of using this composition to screen for analogs thereof, the analogs having anti-HIV activity or HIV transcription repressing activity.

A further object of the present invention is to provide a pharmaceutical composition that represses TRC-associated HIV transcription, thereby providing a novel therapy for HIV infection.

25 In a preferred embodiment, this pharmaceutical composition comprises an effective amount of YY1, LSF and HDAC1, or derivatives or analogs thereof.

A further object of the present invention is to provide a pharmaceutical composition comprising an effective amount of one or more anti-HIV agent, said agent repressing TRC-associated HIV transcription, thereby providing a novel therapy for HIV infection.

30 In another preferred embodiment, this pharmaceutical composition comprises an effective amount of a nucleic acid or combination of nucleic acids comprising: one or more nucleotide sequences encoding YY1 (or a derivative or analog thereof); one or

more nucleotide sequences encoding LSF (or a derivative or analog thereof); and one or more nucleotide sequences encoding HDAC1 (or a derivative or analog thereof).

In another preferred embodiment, this pharmaceutical composition comprises an effective amount of an agent that enhances the binding of a YY1 or a derivative or analog thereof to HDAC1 or a derivative or analog thereof.

In another preferred embodiment, this pharmaceutical composition comprises an effective amount of an agent that enhances recruitment of HDAC1, or a derivative or analog thereof, by YY1, or a derivative or analog thereof, to the HIV LTR RCS site.

In another preferred embodiment, this pharmaceutical composition comprises an effective amount of an agent that enhances the activity of HDAC1, or a derivative or analog thereof, said amount effective to repress HIV transcription.

In another preferred embodiment, this pharmaceutical composition comprises an effective amount of an agent that enhances the expression of HDAC1 or a derivative or analog thereof.

In another preferred embodiment, this pharmaceutical composition comprises an effective amount of an agent that up-regulates the expression of HDAC1 or a derivative or analog thereof.

In another preferred embodiment, this pharmaceutical composition comprises an effective amount of YY1, LSF and HDAC1.

Yet another object of the present invention is to provide methods of identifying, screening, and/or isolating compounds having activities that affect or regulate the TRC complex and its repression of HIV LTR-driven transcription.

A further object of the invention is to provide methods of identifying, screening and/or isolating compounds with "anti-HIV activity". Such compounds find therapeutic utility and may be used to form pharmaceutical compositions for the treatment of active and/or latent HIV infections. Of particular interest are compounds that affect or regulate HDAC1 expression or activity or affect or regulate the HDAC1-recruiting activity of YY1.

In one preferred embodiment, the compounds of interest act to enhance, augment or up-regulate the expression of HDAC1. Such compounds find therapeutic utility in the treatment of active HIV infections.

In another preferred embodiment, the compounds of interest act to enhance or augment the activity of HDAC1. Such compounds find utility in the therapeutic treatment of active HIV infections.

5 In another preferred embodiment, the compounds of interest act to enhance or augment the ability of YY1 to recruit HDAC1 to the RCS site of the HIV LTR. Such compounds find therapeutic utility in the treatment of active HIV infections.

In an alternate embodiment, the compounds of interest act to inhibit, repress or down-regulate the expression of HDAC1. Such compounds find therapeutic utility in the treatment of latent HIV infections.

10 In another embodiment, the compounds of interest act to inhibit or repress the activity of HDAC1. Such compounds find therapeutic utility in the treatment of active HIV infections.

In another preferred embodiment, the compounds of interest act to inhibit or repress the ability of YY1 to recruit HDAC1 to the RCS site of the HIV LTR. Such compounds find therapeutic utility in the treatment of active HIV infections.

15 A further object of the present invention is to provide a method and composition for modulating the histone structure of the RCS site of the HIV LTR so as to modulate the activity of the TRC complex and TRC-associated repression of HIV transcription.

20 Yet another object of the present invention is to provide a method and composition for modulating the association between HDAC1 and YY1 so as to modulate the activity of the TRC complex and TRC-associated repression of HIV transcription.

25 These and other objects, aspects, features, and advantages of the invention will become evident upon reference to the following description.

BRIEF DESCRIPTION OF THE DRAWINGS

Figures 1A-1C depicting the association between YY1 and LSF *in vivo* and *in vitro*, in the absence of a DNA binding site or other factors.

30 Figure 1A is a western blot depicting the immunoprecipitations of Jurkat nuclear extracts using either α -YY1, α -LSF or a nonspecific rabbit polyclonal

antiserum. Mock immunoprecipitations were performed in the absence of antibody. Precipitates were assayed by Western blot using α -LSF. Approximately 75% of the LSF protein recovered by α -LSF is also immunoprecipitated by α -YY1. To demonstrate the recognition of YY1, a western blot of input nuclear extract is displayed at the right.

Figure 1B depicts the EMSA results performed using the RCS binding site and the indicated amounts of LSF and YY1; total amount of protein was normalized by the addition of BSA. The mobility of the native RCS complex formed by nuclear extract is displayed at the right.

Figure 1C depicts the EMSA results performed using the RCS binding site and the indicated amounts of either α -YY1 or α -LSF. Addition of α -YY1 had no effect on the LSF complexes in the absence of YY1 protein (not shown).

Figures 2A-2C map the YY1/LSF interaction domains.

Figure 2A is a representation of LSF deletion mutants used to identify the region of interaction between LSF with YY1. The terminology is defined as follows: ΔA represents a deletion up to codon A, $B\Delta$ represents a deletion after codon B, $A\Delta B$ represents a deletion between codons A and B, XX represents mutated single codons. The amount of LSF bound to GST-YY1 varied from 2.5 to 7% of the input depending on the experiment. All values were normalized to the amount of wild-type LSF bound to GST-YY1 within the experiment.

Figure 2B is a representative autoradiograph showing input LSF constructs, LSF constructs retained by GST-YY1 and by GST, respectively.

Figure 2C is a graphical representation of the YY1 chimeras, all of which contained the GST tag. All constructs also contained the N-terminal region of YY1 (amino acids 1-294) except YZnFs, which lacked this region. YY1 is the full-length wild type YY1 molecule. Non-shaded regions represent GFI-1 zinc fingers (a related Krüppel zinc finger protein). Y/GFI contains only GFI-1 zinc fingers, Chi 1 contained

the first YY1 zinc finger, Chi 2 the first two YY1 zinc fingers, Chi 5 the last two YY1 zinc fingers, Chi 7 the second YY1 zinc finger only, and ZnFs all four YY1 zinc fingers without the YY1 amino-terminal region. The first two zinc fingers of YY1 are required for optimal binding of LSF. Chi 1, Chi 2, Chi 7, and YY1 bound LSF
5 whereas Chi 5, GFI-1 and GST exhibited background levels of LSF binding. A lane containing only a diluted aliquot of labeled LSF serves as a marker. When normalized for protein concentration, a YZnFs construct expressing only the YY1 zinc fingers fused to GST binds LSF with equal avidity to intact GST-YY1. Background levels of binding varied between experiments, as shown.

10 Figure 3 shows that repression by YY1 and LSF require functional LSF and HDAC1 interaction competent YY1. Expression of an integrated LTR-CAT reporter in HeLa-CD4-LTR cells, when activated by 200 ng of pAR-Tat, is inhibited by 2.5 μ g CMV-YY1 or 2.5 μ g of both CMV-YY1 and CMV-LSF. 2.5 μ g CMV-LSF had no effect on expression of CAT. 2.5 μ g of the dominant negative form of LSF (pCMV-
15 LSF 234QL/236KE), incapable of binding DNA but capable of forming inactive multimers, blocks inhibition of Tat-activated LTR expression by 2.5 μ g of YY1. 2.5 μ g CMV-YY1 Δ 154-199, incapable of interacting with HDAC1, was unable to inhibit Tat-activated expression. All transfections received a total of 5 μ g CMV promoter-driven plasmid. Data from at least four independent transfections, normalized for
20 expression of cotransfected β -actin-luciferase.

Figures 4A and 4B show that YY1, LSF and HDAC1 copurify with RCS-binding activity.

Figure 4A shows the activities of crude nuclear extract and elution fractions from an RCS DNA-affinity chromatography column. EMSA (top panel) using the
25 RCS probe. Western blot using rabbit polyclonal α YY1 (second panel), rabbit polyclonal α -CP2 (LSF) antiserum (third panel), and rabbit polyclonal α -HDAC1/2 antibody (bottom panel). EMSA were performed with 4 μ g of nuclear extract and 20

ng of DNA-affinity column eluate. Western blot was performed with 20 µg of nuclear extract and 200ng of DNA-affinity column eluate. Arrow indicates YY1-specific complex, as validated by α-YY1 interference in EMSA. Molecular weight markers are indicated.

5 Figure 4B demonstrates that the HDAC1 activity of DNA affinity chromatography fractions correlates with the presence of the YY1/LSF complex.

Figure 5 is a graph depicting the affect of YY1 on the production of HIV *in vitro*. Production of HIV-1 is inhibited by YY1 but not by YY1Δ154-199 lacking the HDAC1 interaction domain following transfection of HeLa cells with 0.5 µg of the
10 CXCR4 prototypic clone pNL4-3 (left panel), or 1 µg of the CCR5 prototypic clone pYU-2 (right panel). Data is representative of three transfections.

Figure 6 is a model of recruitment by LSF of YY1, and then HDAC1 to the HIV promoter.

Figure 7 is the HIV-1 LTR partial nucleotide sequence (**SEQ ID NO: 1**). Note
15 that the transcription start site is nucleotide +1 (the "g" indicated by arrow); the nucleotides upstream the transcription start site have a negative numeration, while the nucleotides downstream the transcription start site have a positive numeration. The two NF-KB binding sites, the three Sp1 binding sites and the Repressor Complex Sequence (RCS) are labeled.

20 Figure 8 is the human YY1 cDNA nucleotide sequence (**SEQ ID NO: 2**) and amino acid sequence (**SEQ ID NO: 3**)

Figure 9 is the human LSF cDNA nucleotide sequence (**SEQ ID NO: 4**) and amino acid sequence (**SEQ ID NO: 5**)

Figure 10A is the human histone deacetylase 1 amino acid sequence (**SEQ ID**
25 **NO: 6**)

Figure 10B is the human histone deacetylase 1 mRNA nucleotide sequence (**SEQ ID NO: 7**)

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Genetic and biochemical studies have established that chromatin in living cells critically affects transcriptional competence of a promoter sequence (3, 14, 36, 58, 67).

5 A number of recent reports have documented the importance of histone deacetylases as the effector molecules of transcriptional down-regulation in many genes (11, 20, 25, 39, 47). In addition, several transcriptional repressors have been described that tether HDACs to the promoter (2, 3, 21, 28, 31, 42, 43, 71, 72, 74).

We mapped the interactions of LSF and YY1 using a number of chimeric YY1
10 and truncated LSF constructs to determine the domains that participate in complex formation and regulation of the HIV promoter. We subsequently discovered a novel molecular mechanism of repression of an integrated HIV provirus *in vivo*, wherein LSF is required for recruitment of YY1 to the HIV LTR, and repression is mediated by YY1 via the action of HDAC1.

15 We discovered that the YY1/LSF complex copurifies with HDAC1, identified by both Western blot analysis and enzymatic activity assay. Deletion of a glycine/alanine-rich domain of YY1, previously shown to specifically direct the interaction between YY1 and HDAC1 (71), ablates the ability of YY1 to repress the HIV-1 LTR. Further YY1-mediated repression of the LTR is ablated by the
20 deacetylase inhibitor trichostatin A. This is the first discovery of the presence of HDAC1 in the transcription repressor complex. This is also the first discovery of a molecular mechanism by which the TRC represses HIV-1 transcription.

As discussed herein, particularly in the Examples section, we show that LSF and YY1 interact with one another both *in vitro* and *in vivo*. Interaction was observed
25 in the absence of: (a) DNA binding site, (b) other cellular factors, and (c) YY1 C-terminal zinc fingers required for DNA binding to canonical YY1 sites (Figs. 1, 2, and ref. 17). As a majority of the LSF that can be recovered by immunoprecipitation can also be recovered in association with YY1, this complex is likely to be preformed in the cell prior to binding to viral regulatory elements. Further evidence of YY1 and
30 LSF interaction is provided by the observation that YY1 alone does not bind the RCS site in EMSA, but α -YY1 supershifted a significant fraction of the RCS-protein complex.

Overexpression of LSF alone does not repress transcription from the HIV LTR (49, 75). However, an LTR reporter gene is inhibited by YY1 expression, and this effect is augmented by coexpression of LSF. This occurs in the context of both plasmid based (49) and chromatin based reporter genes (**Fig. 3**). Further, both YY1 and LSF function are required for inhibition of the HIV LTR. While expression of LSF does not significantly inhibit LTR expression, LSF synergizes with YY1 in repression, and dominant negative LSF prevents repression by YY1 (**Fig. 3**). Consistent with this model, preliminary studies show that the replication of a provirus containing TAR mutations that block RCS formation, but allow Tat function (48) is unaffected by YY1 (data not shown).

YY1 is known to interact with a number of cellular factors via a Gly/Ala-rich region within residues 154-198 (2). LSF, however, interacts with the zinc finger domain of YY1. The Gly/Ala domain was present on all chimeric YY1 constructs, but only YY1, chimeras 1, 2 and 7, and the YY1 zinc fingers alone were able to specifically bind LSF *in vitro* (**Fig. 2c**). These results indicate that the first and to a lesser extent the second zinc finger domain of YY1 participate in interaction with LSF. While zinc fingers often mediate DNA binding, examples of protein-protein interactions mediated by zinc fingers have been documented (3, 19, 30, 65,).

Our findings show that LSF is required for recruitment of YY1 to the HIV promoter. LSF appears to facilitate YY1 recognition of the LTR, guiding YY1 onto a site that was inaccessible or of low affinity in its absence. However, YY1 may enter the RCS complex solely through protein-protein interaction with LSF. Given the sensitivity of this interaction to mutations within the core of LSF that impair multimerization, it is likely that YY1 recognizes a structure displayed by LSF multimers. The location of the RCS within the HIV LTR may position YY1 to directly inhibit the basal transcription complex or activators of the LTR, as well as recruit mediators of repression such as HDACs.

The multiprotein repressor complex containing the human factors YY1 and LSF, previously isolated from CEM cells, detected in primary T cells, and shown to repress the HIV-1 LTR, copurifies with a 65-kDa protein which we have identified as HDAC1 (**Fig. 4a**). The anti-HDAC1 antibody used to perform the Western blot analysis was reactive to both HDAC1 and HDAC2, but not to HDAC3 (72).

Molecular weight found in western blot analysis (59, 71) suggest that HDAC1 is the protein copurified with YY1 and LSF.

We have demonstrated that the Gly/Ala-rich domain of YY1, which mediates the interaction with HDAC1 (71), is required for efficient repression of the HIV-1 LTR by the YY1-LSF complex (**Fig. 3**). This suggests that recruitment of a histone deacetylase is a necessary event in the mechanism of repression of HIV-1 gene expression by YY1. Indeed, the fact that transfection of YY1 Δ 154-199 resulted in modest up-regulation of LTR expression and HIV production suggests that LTR expression may in part reflect the competing influences of cellular HDAC1 and histone acetyltransferase activity.

Studies of the chromatin structure of the integrated HIV-1 provirus in chronically and acutely infected cells lines have detected the presence of a large nucleosome-free, DNase I-hypersensitive region spanning nucleotides 223 to 450 of the HIV-1 genome. This corresponds to the portion of the LTR including the enhancer and the promoter regions, up to the transcription start site. Upon treatment with TPA or TNF- α , the 3' boundary of the nucleosome-free region was extended a further 140 nucleotides, indicating the alteration of the nucleosome, termed nuc-1 (51, 61-63). Additional DNase I-hypersensitive sites and nucleosome-protected regions have been identified all along the integrated HIV-1 genome (61, 62).

More recently Pazin et al (46) have shown that binding of both Sp1 and the p50 subunit of NF-KB to the HIV-1 LTR alters the local nucleosomal array in vicinity of the HIV-1 promoter and produces the DNase I-hypersensitive region between nucleotides 223 and 450. However, it is the p65 subunit of NF-KB that induces changes in the nucleosome nuc-1, perhaps through the recruitment of a histone acetyltransferase (51), and enhances transcriptional activity (46).

Our previous results suggested that LSF allowed YY1 to recognize a site on the LTR that YY1 could not bind by itself (49). Therefore, LSF might primarily act as a docking molecule for YY1, which in turn acts by tethering histone deacetylase (**Fig. 6**). In this model YY1 may be a limiting factor for repression of the LTR, required for the recruitment of the histone deacetylase to the HIV-1 promoter. The finding that overexpression of the mutant YY1 Δ 154-199 results in activation of HIV expression is consistent with such a model. El Kharroubi et al (13) has shown that activation of the

integrated proviral HIV genome requires alteration of the local chromatin via acetylation of the nucleosome adjacent to the start site. The recruitment of HDAC1 by YY1 might prevent such changes in the local chromatin, maintaining the nucleosome in a deacetylated state, preserving higher order nucleosome structure and thereby inhibiting gene expression (**Fig. 6**).

In vitro assays have shown that assembly of nuc-1 on naked HIV-1 DNA can be inhibited by the presence of LSF (51). Further, one previous report suggested that LSF is important for efficient activation of the HIV-1 LTR (26), although this has been disputed (75). However, we find no evidence that wild type LSF activates LTR expression. While previous data that LSF is an activator of HIV conflicts with our findings, these studies were performed in very different experimental systems. Indeed, through interaction with other factors, in the absence of YY1, or in other cellular milieus LSF might direct LTR activation.

The HIV-1 enhancer and promoter possess a multiplicity of sequences recognized by cellular and viral regulatory factors. The role of cellular enhancers such as Sp1 and NF-KB, and the viral activator Tat in active HIV gene expression has been extensively studied. As discussed above, changes in chromatin structure about an integrated HIV promoter during activation have been documented. However, mechanisms that down-regulate HIV expression are largely unknown. We have shown that YY1 and LSF are capable of cooperating to inhibit HIV transcription. Of the many possible mechanisms through which YY1 might down-regulate transcription, we can now link this function to the recruitment of a histone deacetylase; our studies strongly suggest this enzyme is histone deacetylase 1. Thus the YY1-LSF repressor complex recruits factors capable of potent and durable inhibition of HIV-1 LTR promoter expression.

A large nucleosome-free, DNase I-hypersensitive region spanning nucleotides 223 to 450 of the HIV-1 has been observed in the chromatin structure of the integrated HIV-1 provirus in chronically and acutely infected cells lines. Activation of LTR expression extends the 3' boundary of this nucleosome-free region a further 140 nucleotides. Each nucleosome is entwined by 1.65 turns of a left-handed superhelix of DNA that corresponds to 147 basepair. This indicates that the DNA protected by one nucleosome has been exposed, presumably by remodeling of the nucleosome structure.

The binding of both Sp1 and the p50 or p65 subunits of NF- κ B to the HIV-1 LTR alters the local nucleosomal array in vicinity of the HIV-1 promoter, and perhaps through the recruitment of a histone acetyltransferase enhances transcriptional activity. El Kharroubi *et al* (13) have also shown that activation of the integrated proviral HIV genome requires alteration of the local chromatin via acetylation of the nucleosome adjacent to the start site. Our findings imply that recruitment of HDAC1 by YY1 might prevent such changes in the local chromatin, maintaining the nucleosome in a deacetylated state and inhibiting HIV expression.

We propose a dynamic model of HIV LTR regulation that would allow the establishment of virological latency in rare CD4 T cells. Following T cell activation necessary for viral entry, reverse transcription, and other steps of the viral life cycle which lead to proviral integration, the rare activated cell avoids apoptosis or viral or immune-mediated destruction. Dampening of LTR expression by YY1 and LSF may play an important role at this stage. This cell then follows pathways that typically reestablish the resting, memory state. The HIV-1 LTR remains silent due to the predominant effects of repressor molecules, resulting in an inaccessible chromatin structure about the LTR. This cell may later exit virological latency if it encounters appropriate stimuli that to increase nuclear levels of NF- κ B, again changing LTR chromatin structure. Levels of the viral activator tat then increase within the cell, driving the equilibrium towards viral expression.

Definitions:

As used herein, the "HIV LTR" refers to the long term repeat sequence on the HIV that acts as the primary promoter of HIV transcription. The repressor complex sequence or "RCS" refers to the region -10 to +27 of the HIV-1 LTR. The HIV LTR sequence is set forth in Figure 8 (**SEQ ID NO: 1**).

As used herein, the transcription repressor complex or "TRC" refers to a protein complex containing YY1, LSF, and HDAC1 that binds to the LTR RCS and inhibits HIV transcription.

As used herein, Yin Yang 1 or "YY1" (also termed δ , NF-E1, UCRBP or CF1; refs. 45, 52, 56, 40, 69) is a cellular transcription factor that has been shown to bind to

the RCS and inhibit LTR-driven HIV transcription. The protein and cDNA sequences for YY1 are set forth in Figure 8 (**SEQ ID NOS: 2 and 3**).

As used herein, "LSF" (also termed LBF-1, CP-2, LBP-1a, b, c and d) is a lymphoid transcription factor that has been shown to repress LTR-driven transcription.

5 The protein and cDNA sequences for LSF are set forth in Figure 9 (**SEQ ID NOS: 4 and 5**).

As used herein, "HDAC1" refers to histone deacetylase 1, an enzyme that hydrolyzes n-acetyl groups on histones, nucleic proteins found in most eukaryotic cells. Histones are generally complexed to DNA in chromatin and chromosomes
10 responsible for compacting DNA enough so that it will fit within a nucleus. Histones are generally of relatively low molecular weight and are basic, having a very high arginine/lysine content. They are highly conserved and may act as nonspecific repressors of gene transcription. The protein and mRNA sequences for HDAC1 are set forth in Figures 10A and 10B (**SEQ ID NOS: 6 and 7**).

15

Proteins, Peptides, Derivative and Analogs:

The invention provides compositions comprising or, alternatively, consisting of or consisting essentially of, an isolated transcription repressor complex, comprised of YY1 and LSF, and HDAC1 proteins.

20 As used herein, a peptide is said to be "isolated" or "purified" when it is substantially free of homologous cellular material or chemical precursors or other chemicals. The peptides of the present invention can be purified to homogeneity or other degrees of purity. The level of purification will be based on the intended use.

The composition of the invention may contain a derivative (*e.g.*, a fragment) or
25 analog of YY1, LSF, and/or HDAC1.

As used herein, the terms "derivative" and "analog" refer proteins, the amino acid sequence of which consists of a portion the full length protein, which portion retains the activity of the full length protein. Activity or effectiveness of the proteins, derivatives and/or analogs of the invention for treatment or prevention of HIV
30 infection can be determined by any of the methods by known in the art, disclosed herein and/or described in parent patent application, U.S. Serial No. 09/355,010 and

International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

In the context of the instant invention, derivatives and/or analogs of the YY1, LSF and/or HDAC1 proteins retain activity sufficient to form the transcription repressor complex, bind to the RCS site of HIV LTR, and/or inhibit HIV transcription. In a preferred embodiment, the composition of the invention contains a YY1, LSF and/or HDAC1 derivative, the amino acid sequence of which consists of one or more functional domains of the YY1, LSF and/or HDAC1 proteins. In various specific embodiments, the portion of the YY1, LSF, and/or HDAC1 sequence is at least 10, 20, 30, 40, 50, 60, 80, 100, 200, 300, or 400 amino acids.

As used herein, a "functional domain" is the region of a protein (i.e., amino acid sequence) or an oligonucleotide (e.g., nucleic acid sequence) that is primarily responsible for the function of the protein (e.g., catalytic site of an enzyme, binding epitope of an antibody, etc.), the absence of which would result in loss of function.

The functional domain of YY1 has been analyzed and described by Bushmeyer *et al.* (Bushmeyer *et al.*, 1995, J. Biol. Chem. 270:30213). The transcriptional repression domain was mapped in the region between amino acids 333 and 397 of the YY1 amino acid sequence (SEQ ID NO:2); therefore, it overlaps with the three last zinc finger domains. A more detailed mapping of the repression domain has been obtained in Yang Shi's laboratory. Interaction with other transcription factors has been shown to alter and regulate YY1 action (Seto *et al.*, 1993, Nature 365:462; Lee *et al.*, 1993, Proc. Natl. Acad. Sci. USA 90:6145; Lee *et al.*, 1995, Nucleic Acid Research 23:925; Lee *et al.*, 1995, Genes & Development 9:1188; Lewis *et al.*, 1995, J. Virology 69:1628; Inouye and Seto, 1994, J. Biol. Chem. 269:6506). Several regions of YY1 have been shown to be involved in protein-protein interaction with transcription factors that regulate YY1 action: amino acids 260-331 are required for interaction with Sp1; amino acids 201-343 for interaction with c-Myc; amino acids 332-414 for interaction with E1A; and amino acids 224-330 and 332-414 are necessary for binding to ATF-2a (Bushmeyer *et al.*, 1995, J. Biol. Chem. 270:30213; Zhou *et al.*, 1995, J. Virology 69:4323). Further analysis of the LSF functional domain is provided in the Examples section below.

Preferred YY1 derivatives and analogs comprise the following sequences:
amino acid numbers 50-414, 101-414, 150-414, 175-414, 200-414, 250-414, 260-414,
270-414, 280-414, 290-414, 300-414, 320-414, 340-414 and 360-414, and, most
preferably, amino acid numbers 200-414, of the YY1 sequence as depicted in Figure 8
5 (SEQ ID NO:3).

The LSF functional domains have been examined by Shirra et al. (Shirra *et al.*,
1994, Molecular and Cellular Biology 14:5076). LSF binds the DNA as a tetramer.
The region between amino acids 189 and 239 of the LSF amino acid sequence (SEQ
ID NO:5) appears to be necessary for binding to the DNA. Further analysis of the
10 LSF functional domain is provided in the Examples section below. For example, our
studies show that truncations upstream of LSF amino acid 183 tend to disrupt the
interaction between LSF and YY1. Likewise, interaction with YY1 appears to involve
LSF amino acids 164-403.

In the context of the instant invention, preferred LSF derivatives and analogs
15 containing the following amino acid sequences: amino acid numbers 150-250 and 200-
300, and, most preferably, amino acid numbers 189-239 of the LSF sequence as
depicted in Figure 9 (SEQ ID NO:5). Preferred fragments are those less than 75, 100,
150, 200, 250, or 300 amino acids in length.

The HDAC1 functional domains relevant to its recruitment by YY1 are
20 described herein, particularly in the Examples section, as well as in by WM Yang (71,
72).

In one embodiment, derivatives can be made by altering the amino acid
sequence of the protein by substitutions, additions or deletions that provide for
therapeutically effective molecules. Thus, the derivatives include peptides containing,
25 as a primary amino acid sequence, all or part of the HDAC1, YY1 and/or LSF amino
acid sequence including altered sequences in which functionally equivalent amino acid
residues are substituted for residues within the sequence resulting in a peptide which is
functionally active.

For example, one or more amino acid residues within the sequence can be
30 substituted by another amino acid of a similar polarity which acts as a functional
equivalent, resulting in a silent alteration. Conservative substitutions for an amino
acid within the sequence may be selected from other members of the class to which the

amino acid belongs. For example, the nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan and methionine. The polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, asparagine, and glutamine. The positively charged (basic) amino acids include arginine, lysine and histidine. The negatively charged (acidic) amino acids include aspartic acid and glutamic acid. Any technique for mutagenesis known in the art can be used, including but not limited to, chemical mutagenesis, *in vitro* site-directed mutagenesis (Hutchinson *et al.*, 1978, J. Biol. Chem 253:6551), use of TAB7 linkers (Pharmacia), PCR with primers containing mutations, etc.

In certain embodiments, it is desirable to introduce nonclassical amino acids or chemical amino acid analogs as a substitution or addition into the HDAC1, YY1 and LSF proteins and/or derivatives. Non-classical amino acids include but are not limited to the D-isomers of the common amino acids, 2,4-diaminobutyric acid, α -amino isobutyric acid, 4-aminobutyric acid, Abu, 2-amino butyric acid, γ -Abu, ϵ -Ahx, 6-amino hexanoic acid, Aib, 2-amino isobutyric acid, 3-amino propionic acid, ornithine, norleucine, norvaline, hydroxyproline, sarcosine, citrulline, homocitrulline, cysteic acid, t-butylglycine, t-butylalanine, phenylglycine, cyclohexylalanine, β -alanine, fluoro-amino acids, designer amino acids such as β -methyl amino acids, α -methyl amino acids, $N\alpha$ -methyl amino acids, and amino acid analogs in general. Furthermore, the amino acid can be D (dextrorotary) or L (levorotary).

The invention also provides HDAC1, YY1 and LSF, derivatives or analogs that are cyclized and/or branched using techniques known in the art

Also included within the scope of the invention are HDAC1, YY1 and LSF, or derivatives or analogs thereof, which are differentially modified during or after synthesis, *e.g.*, by benzylation, glycosylation, acetylation, phosphorylation, amidation, pegylation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to an antibody molecule or other cellular ligand, etc. In specific embodiments, the serine residues of YY1 and LSF, or derivatives or analogs thereof, are phosphorylated using techniques known in the art. In other specific embodiments, YY1 and LSF, or derivatives or analogs thereof, are acetylated at the N-terminus and/or amidated at the C-terminus. Any of numerous chemical modifications may be

carried out by known techniques, including but not limited to acetylation, formylation, oxidation, reduction; metabolic synthesis in the presence of tunicamycin; etc.

The derivative and/or analog may further comprise a chimeric, or fusion, protein comprising a component of the TRC (e.g., YY1, LSF, or HDAC1) or a functional derivative or analog thereof joined at its amino- or carboxy-terminus via a peptide bond to an amino acid sequence of another HIV transcription factor, preferably one of the proteins of interest mentioned above. In a specific embodiment, the chimeric or fusion protein comprises an at least six amino acid portion, or an at least 10, 20, 30, 40, 50, 75, 100 or 200 amino acid portion, of one protein of interest joined via a peptide bond to an at least six amino acid portion, or an at least 10, 20, 30, 40, 50, 75, 100 or 200 amino acid portion, of another protein of interest, preferably where said portions are active to treat or prevent HIV infection. Exemplary procedures for producing or expression such chimeric proteins are known in the art and/or described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

HDAC1, YY1 and LSF derivatives and analogs can be made by chemical synthesis or by recombinant production from nucleic acid encoding HDAC1, YY1 and LSF peptide which nucleic acid has been mutated. Exemplary methods for chemically or recombinantly synthesizing proteins, derivatives and analogs are known in the art and/or fully described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

Nucleic Acids and Oligonucleotides:

The invention further provides nucleic acids comprising nucleotide sequences encoding YY1, or derivatives, fragments or analogs thereof; LSF, or derivatives fragments or analogs thereof; and HDAC1, or derivatives fragments or analogs thereof.

The nucleotide sequences encoding, and the corresponding amino acid sequences of, HDAC1, LSF and YY1 are known (Shi *et al.*, 1991, Cell 67:377-388 and Kato *et al.*, 1991, Science 251:1476, respectively) and are provided in Figures 8-10,

respectively (SEQ ID NOS:2-7, respectively). Nucleic acids encoding HDAC1, LSF and YY1 can be obtained by any method known in the art, *e.g.*, by PCR amplification using synthetic primers hybridizable to the 3' and 5' ends of the sequence and/or by cloning from a cDNA or genomic library using an oligonucleotide specific for the gene sequence.

Methodologies for identifying, isolating, and amplifying nucleotide sequences associated with HDAC1, LSF and YY1 are known in the art and/or fully described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

The HDAC1, LSF or YY1 sequences provided by the instant invention include those nucleotide sequences encoding substantially the same amino acid sequences as found in native proteins, and those encoded amino acid sequences with functionally equivalent amino acids, as well as those encoding other derivatives or analogs.

Homologs (*e.g.*, nucleic acids encoding HDAC1, LSF and YY1 of species other than human) or other related sequences (*e.g.*, paralogs) can also be obtained by low, moderate or high stringency hybridization with all or a portion of the particular human sequence as a probe using methods well known in the art for nucleic acid hybridization and cloning. Exemplary hybridization procedures are known in the art and/or fully described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

Inhibitors, Antibodies, and Antisense:

The invention further provides for inhibitors of YY1, LSF and HDAC1 (or derivatives or analogs thereof), particularly those agents that (a) inhibit transcription repressor complex formation (*i.e.*, the interaction between YY1, LSF and HDAC1); (b) inhibit the activity of one or more of the TRC components; (c) inhibit the binding of the TRC to the HIV LTR; and/or (d) prevent TRC-associated repression of HIV transcription. Such inhibitors may be identified by any method known in the art for assaying formation of the TRC; interaction among the TRC components; binding of the TRC to the HIV LTR or to its mediators; and/or HIV transcription, infection, or replication. Exemplary methods are known in the art, described herein and/or

described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

5 The compounds that may be screened in accordance with the invention include, but are not limited to, peptides, antibodies and fragments thereof, and other organic compounds (*e.g.*, peptidomimetics) that inhibit formation of the TRC, activity of the TRC, or binding of the TRC to the LTR of HIV. These screens identify peptides, antibodies or fragments thereof, and other organic compounds that inhibit suppression of HIV transcription mediated by the TRC.

10 Such compounds may include, but are not limited to, peptides such as, for example, soluble peptides, including but not limited to, those found in random peptide libraries; (see, *e.g.*, Lam *et al.*, 1991, Nature 354:82-84; Houghten *et al.*, 1991, Nature 354:84-86). Such compounds may also be found in combinatorial chemistry-derived molecular libraries made of D- and/or L- configuration amino acids; phosphopeptides
15 (including, but not limited to, members of random or partially degenerate, directed phosphopeptide libraries; see, *e.g.*, Songyang *et al.*, 1993, Cell 72:767-778); antibodies (including, but not limited to, polyclonal, monoclonal, humanized, anti-idiotypic, chimeric or single chain antibodies, and FAb, F(abN)₂ and FAb expression library fragments, and epitope-binding fragments thereof); antisense RNA and small organic
20 or inorganic molecules. In a specific embodiment, pyrrole-imidazole polyamides (*e.g.* as described in Gottesfeld *et al.*, 1997, Nature 387:202-205) are provided to inhibit the activity of the transcription repressor complex on HIV gene expression.

By way of examples of non-peptide libraries, a benzodiazepine library (see
25 *e.g.*, Bunin *et al.*, 1994, Proc. Natl. Acad. Sci. USA 91:4708-4712) can be adapted for use. Peptoid libraries (Simon *et al.*, 1992, Proc. Natl. Acad. Sci. USA 89:9367-9371) can also be used. Another example of a library that can be used, in which the amide functionalities in peptides have been permethylated to generate a chemically transformed combinatorial library, is described by Ostresh *et al.* (1994, Proc. Natl. Acad. Sci. USA 91:11138-11142).

In another embodiment of the invention, the TRC, protein components thereof, transcription mediators recruited thereby, or fragments, derivatives, or analogs these proteins, may be used as an immunogen to generate antibodies that recognize such an immunogen. Such antibodies include but are not limited to polyclonal, monoclonal, 5 chimeric, single chain, Fab fragments, and an Fab expression library. In one embodiment, antibodies that specifically bind to YY1 or LSF and prevent TRC formation are provided. In another embodiment, antibodies that bind the TRC and prevent its binding to the HIV LTR are provided.

Various procedures known in the art may be used for the production of 10 polyclonal antibodies and monoclonal antibodies to HDAC1, LSF, YY1, and/or the transcription repressor complex, or derivative or analog thereof. Exemplary procedures are described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

15 Antibody fragments and other derivatives which contain the idiotype (binding domain) of the molecule can be generated by known techniques. For example, such fragments include but are not limited to: the F(abN)₂ fragment which can be produced by pepsin digestion of the antibody molecule; the FabN fragments which can be generated by reducing the disulfide bridges of the F(abN)₂ fragment; and the Fab 20 fragments which can be generated by treating the antibody molecule with papain and a reducing agent.

The function of the TRC or the individual protein components thereof (e.g., LSF, YY1, and HDAC1) can be inhibited by use of antisense nucleic acids for HDAC1, LSF and/or YY1. The present invention provides the therapeutic or 25 prophylactic use of nucleic acids of at least six nucleotides that are antisense to a gene or cDNA encoding HDAC1, LSF and/or YY1, or portions thereof.

An "antisense" nucleic acid as used herein refers to a nucleic acid capable of hybridizing to a sequence-specific (e.g., non-poly A) portion of the protein RNA (preferably mRNA) by virtue of some sequence complementarity. The antisense 30 nucleic acid may be complementary to a coding and/or noncoding region of the protein mRNA. Such antisense nucleic acids have utility as therapeutics that inhibit transcription repressor complex formation or activity, or HDAC1, LSF, or YY1

function or activity, and can be used in the treatment or prevention of disorders as described *supra*.

The antisense nucleic acids of the invention can be oligonucleotides that are double-stranded or single-stranded, RNA or DNA or a modification or derivative thereof, which can be directly administered to a cell, or which can be produced
5 intracellularly by transcription of exogenous, introduced sequences.

The antisense nucleic acids of the present invention are preferably at least six nucleotides and are more preferably oligonucleotides (ranging from 6 to about 200 oligonucleotides). In specific aspects, the oligonucleotide is at least 10 nucleotides, at
10 least 15 nucleotides, at least 100 nucleotides, or at least 200 nucleotides.

The oligonucleotides can be DNA or RNA or chimeric mixtures or derivatives or modified versions thereof, single-stranded or double-stranded. In a preferred embodiment, the antisense oligonucleotide is a single-stranded DNA.

The oligonucleotide may be modified at any position on its structure with
15 constituents generally known in the art. The oligonucleotide can be modified at the base moiety, sugar moiety, or phosphate backbone. The oligonucleotide may include other appending groups such as peptides, or agents facilitating transport across the cell membrane (see, *e.g.*, Letsinger et al., 1989, Proc. Natl. Acad. Sci. U.S.A. 86: 6553-6556; Lemaitre et al., 1987, Proc. Natl. Acad. Sci. 84: 648-652; PCT Publication
20 No. WO 88/09810, published December 15, 1988) or blood-brain barrier (see, *e.g.*, PCT Publication No. WO 89/10134, published April 25, 1988), hybridization-triggered cleavage agents (see, *e.g.*, Krol et al., 1988, BioTechniques 6: 958-976) or intercalating agents (see, *e.g.*, Zon, 1988, Pharm. Res. 5: 539-549). Exemplary modified moieties suitable for use with the antisense oligonucleotides of the present
25 invention are known in the art and/or described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

The oligonucleotide may be conjugated to another molecule, *e.g.*, a peptide, hybridization triggered cross-linking agent, transport agent, hybridization-triggered
30 cleavage agent, etc.

Oligonucleotides of the invention may be synthesized by standard methods known in the art. Exemplary synthesis methods are described in parent patent

application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

The antisense oligonucleotides comprise catalytic RNAs, or ribozymes (see, e.g., PCT International Publication WO 90/11364, published October 4, 1990; Sarver et al., 1990, Science 247: 1222-1225). In another embodiment, the oligonucleotide is a 2N-0-methylribonucleotide (Inoue et al., 1987, Nucl. Acids Res. 15: 6131-6148), or a chimeric RNA-DNA analog (Inoue et al., 1987, FEBS Lett. 215: 327-330).

The antisense nucleic acids of the invention comprise a sequence complementary to at least a portion of an RNA transcript of the HDAC1, YY1, and/or LSF genes, preferably a human genes. However, absolute complementarity, although preferred, is not required. A sequence "complementary to at least a portion of an RNA," as referred to herein, means a sequence having sufficient complementarity to be able to hybridize with the RNA, forming a stable duplex; in the case of double-stranded antisense nucleic acids, a single strand of the duplex DNA may thus be tested, or triplex formation may be assayed. The ability to hybridize will depend on both the degree of complementarity and the length of the antisense nucleic acid. One skilled in the art can ascertain a tolerable degree of mismatch by use of standard procedures to determine the melting point of the hybridized complex.

In another embodiment, the invention is directed to methods for inhibiting the expression of HDAC1, LSF and/or YY1 nucleic acid sequences in a cell comprising providing the cell with an effective amount of a composition comprising an antisense nucleic acid of LSF, YY1 and/or HDAC1, or derivatives thereof, of the invention. Cell types that express HDAC1, LSF or YY1 RNA can be identified by various methods known in the art. Such methods include, but are not limited to, hybridization with HDAC1, LSF and YY1-specific nucleic acids (e.g. by northern hybridization, dot blot hybridization, in situ hybridization), or by observing the ability of RNA from the cell type to be translated *in vitro* into HDAC1, LSF and YY1 by immunohistochemistry. In a preferred aspect, primary tissue from a patient can be assayed for HDAC1, LSF and/or YY1 expression prior to treatment, e.g., by immunocytochemistry or in situ hybridization.

Enhancers, Activators and Regulators

The invention further provides for agents that affect or regulate the TRC-associated mechanism of repression. Techniques and procedures for identifying and screening for enhancers and regulators are analogous to those used to identify inhibitors, which are described and exemplified in the previous section.

As mentioned above, the invention provides for agents that inhibit, repress or down-regulate the TRC-associated mechanism of repression. Examples of such agents include: agents that repress or inhibit the formation of the TRC; agents that repress or inhibit the binding of YY1 (or a derivative or analog thereof) to HDAC1 (or a derivative or analog thereof); agents that repress or inhibit recruitment of HDAC1 (or a derivative or analog thereof) by YY1 (or a derivative or analog thereof) to the HIV LTR RCS site; agents that repress or inhibit the activity of HDAC1 (or a derivative or analog thereof); agents that repress or inhibit the expression of HDAC1 (or a derivative or analog thereof); and agents that down-regulate the expression of HDAC1 (or a derivative or analog thereof). These agents find particular therapeutic utility in blocking the repression of LTR-driven HIV transcription so as to treat or prevent of latent HIV infection.

In one embodiment, the regulator may comprise an inhibitor of the enzymatic activity of HDAC1. Exemplary HDAC1 inhibitors include trichostatin-A and trapoxin (Hassig CA *et al.*, 1998, Proc Natl Acad Sci U S A 95(7):3519-24).

In another embodiment, the regulator may comprise an agent that modulates expression and/or activity of HDAC1. For example, a dynamic balance of histone acetylation/deacetylation is maintained by histone acetyltransferases and histone deacetylases (Hu, E *et al.*, 2000, J Biol Chem 275(20):15254-64).

In another embodiment, the regulator may comprise an agent that provides a post-translational modification (e.g., phosphorylation) to HDAC1, YY1 and/or LSF.

The invention further provides for agents that enhance, activate or up-regulate the TRC-associated mechanism of repression. Examples of such agents include: agents that enhance the formation of the TRC; agents that enhance the binding of YY1 (or a derivative or analog thereof) to HDAC1 (or a derivative or analog thereof); agents that enhance recruitment of HDAC1 (or a derivative or analog thereof) by YY1 (or a derivative or analog thereof) to the HIV LTR RCS site; agents that enhance the

activity of HDAC1 (or a derivative or analog thereof); agents that enhance the expression of HDAC1 (or a derivative or analog thereof); and agents that up-regulate the expression of HDAC1 (or a derivative or analog thereof). These agents find particular therapeutic utility in the repression of LTR-driven HIV transcription and the treatment or prevention of active HIV infection.

Enhancers and regulators, like inhibitors, may be identified by any method known in the art for assaying for formation of the TRC; interaction among the components of the TRC; binding of the TRC to the HIV LTR or to its mediators; and/or HIV transcription, infection, or replication. Exemplary methods are known in the art, described herein and/or described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein. For example, a standard ELISA assay using an anti-HDAC1 antibody (such as that described in the Examples section below) as substrate may be used to measure the level of HDAC1 expressed in the presence (test) and absence (control) of a particular compound. A measured increase in expression in the presence of a compound is indicative of a compound's ability to enhance or up-regulate the expression of HDAC1. Likewise, co-precipitation assays (such as those described in the Examples section) using HDAC1 and YY1, for example, may be used to measure the binding affinity between the two molecules.

Assays:

Testing for Anti-HIV Activity

Anti-HIV activity of a therapeutic preparation can be measured by assaying it for the ability to inhibit HIV replication, transcription or expression of HIV RNA or protein. Any assay for HIV expression, replication or transcription, either *in vivo* or *in vitro*, can be used to determine the level of transcription repression. For example, but not by way of limitation, EMSA for binding to the HIV LTR, the viral infection assays, CAT or other reporter gene transcription assays (with the CAT reporter gene or any other reporter gene known in the art operably linked to the HIV LTR), HIV infection assays, or assays for viral production from cells latently infected with HIV (for example, but not limited to, by the method described by Chun et al., 1977, Nature 387:183-188) can be used to screen for and test potential inhibitors of YY1-LSF

complexes. Additional methods for assaying the efficacy of a particular preparation or therapy for HIV transcription repression are known in the art and/or described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

5 The therapeutics of the invention are preferably tested *in vitro*, and then *in vivo* for the desired therapeutic or prophylactic activity, prior to use in humans. Any *in vitro* or *in vivo* assay known in the art to measure HIV infection, production, replication or transcription can be used to test the efficacy of a therapeutic of the invention. Exemplary *in vitro* or *in vivo* assays described herein as well as in parent
10 patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

 In vivo animal models for testing the efficacy of the therapeutic preparation of the present invention are known in the art. Once the therapeutic preparation has been tested *in vitro*, and also preferably in a non-human animal model, the utility of the
15 therapeutic preparation can be determined in human subjects. Exemplary human and animal assays are described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

20 Formation and Binding of Transcription Repressor Complex

 TRC formation can be assayed by various methods, including but not limited to, protein affinity chromatography, affinity blotting, immunoprecipitation, cross-linking, and competitive inhibition assay methods (see generally, Phizicky *et al.*, 1995, Microbiol. Rev. 59:94-123). Additionally, since the DNA encoding HDAC1, YY1
25 and LSF have been isolated and sequenced (see *e.g.*, Shi *et al.*, 1991, Cell 67:377-388 and Kato *et al.*, 1991, Science 251:1476, respectively), this sequence may be routinely manipulated in known assays, to identify derivatives, fragments and analogs that bind counterpart members of the HIV-1 LTR binding, repressor complex. Such assays include, but are not limited to, *in vitro* cell aggregation and interaction trap assays (see
30 generally, Phizicky *et al.*, 1995, Microbiol. Rev. 59:94-123).

 The affinity of derivatives and analogs for counterpart members of the repressor complex can routinely be determined by, for example, competitive inhibition

experiments using HDAC1, YY1 and LSF, respectively. In specific embodiments, the derivatives or analogs of the invention display an affinity for the counterpart HIV-1 LTR binding complex member, which affinity approximates or is greater than the affinity of the protein from which it is derived.

5 The ability of complexes comprising YY1 (or derivatives and analogs thereof); LSF (or derivatives and analogs thereof); and HDAC1 (or derivatives and analogs thereof) to bind to the LTR of HIV-1 may routinely be determined using known assays, such as, for example, footprint and electrophoretic mobility shift assays (*e.g.*, see Section 7, *infra*). These assays may routinely be applied to ascertain the affinity of
10 the complex for DNA sequences of the LTR. In one preferred embodiment, the compositions of the invention containing YY1, LSF, and HDAC1 or derivatives and analogs thereof form complexes having the highest affinity for the DNA sequence of the HIV-1 LTR. In a further preferred embodiment, the compositions of the invention are form complexes that bind the DNA sequence corresponding to nucleotides -17 to
15 +17 of the HIV-1 LTR as depicted in Figure 7 (SEQ ID NO:1).

Transcriptional repression of HIV-1 by HDAC1, and derivatives and analogs thereof; LSF, and derivatives and analogs thereof; and YY1, and derivatives and analogs thereof, may routinely be examined using known techniques, such as, for may routinely be examined using known techniques, such as, for example, *in vitro*
20 transcription experiments in which the HIV-1 LTR is operably linked to a reporter gene, such as, for example and not by way of limitation chloramphenicol acetyltransferase (CAT) (see *e.g.*, Section 6 and 7, *infra*).

Assays for binding to the HIV LTR (*e.g.*, electrophoretic mobility shift assay or EMSA) are also useful for testing the efficacy of therapeutic preparations of the
25 invention. Specifically, the therapeutic preparations to be tested is incubated with radioactively labeled, double-stranded DNA containing the nucleotide sequence of -17 to +27 or -10 to +27 of the HIV LTR sequence and then analyzed by non-denaturing gel electrophoresis. A shift in the mobility of the labeled HIV LTR probe after incubation with the therapeutic preparation to be tested indicates that it binds to the
30 HIV LTR.

Screening Assays

Screening of compounds or compound libraries of putative inhibitors can be accomplished by any of a variety of commonly known methods described herein or in parent patent application, U.S. Serial No. 09/355,010 and International Publication No.

5 WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

In a specific embodiment, screening can be carried out by contacting a single compound or multiple library members with an HDAC1, LSF or YY1 protein or derivative, or a repressor complex, or an HIV LTR nucleic acid immobilized on a solid phase and harvesting those library members that bind to the protein (or complex or
10 nucleic acid or derivative).

In a specific embodiment, fragments and/or analogs of YY1 or LSF, especially peptidomimetics, are screened for activity as competitive or non-competitive inhibitors of YY1-LSF complex formation or binding of the complex to the HIV LTR, and thereby for the ability to inhibit YY1-LSF complex activity.

15 Numerous experimental methods may be used to select and detect proteins or non-protein molecules that interfere with the formation of the repressor complex or binding to the HIV LTR and thereby modulate HIV transcription including, but not limited to, protein affinity chromatography, affinity blotting, immunoprecipitation, cross-linking, and library based methods such as protein probing, phage display and
20 the two-hybrid system. See generally, Phizicky *et al.*, 1995, Microbiol. Rev. 59:94-123. For example, the two-hybrid system may be used to detect inhibitors of the interaction between LSF and YY1 by constructing the appropriate hybrids and testing for reporter gene activity in the presence of candidate inhibitors.

Any assay for HIV infection, replication or transcription, either *in vivo* or *in*
25 *vitro*, can be used to screen for inhibitors of the YY1-LSF complex activity.

In the production of antibodies, screening for the desired antibody can be accomplished by techniques known in the art, *e.g.* ELISA (enzyme-linked immunosorbent assay). For example, to select antibodies that recognize a specific domain of LSF, YY1 or a YY1-LSF complex, one may assay generated hybridomas
30 for a product that binds to a fragment containing such domain. For selection of an antibody specific to human LSF, YY1 or YY1-LSF complex, one can select on the

basis of positive binding to a human protein or complex and a lack of binding to the protein or complex of another species, *e.g.* mouse, rat, primate, etc.

Therapeutic Utilities:

5 The invention provides for repression of HIV transcription to treat diseases and disorders associated with HIV infection by administration of a therapeutic compound (termed herein "therapeutic"). Such "therapeutics" include, but are not limited to: compositions containing YY1, LSF, and HDAC1 and therapeutically and prophylactically effective derivatives (including fragments) and/or analogs thereof,
10 *i.e.*, those derivatives and/or analogs which prevent or treat HIV infection (*e.g.*, as demonstrated in *in vitro* and *in vivo* assays described *infra*), as well as nucleic acids encoding YY1, LSF, and HDAC1, and/or therapeutically and prophylactically effective derivatives and analogs thereof (*e.g.*, for use in gene therapy); modulators (*e.g.*, antagonists, inhibitors and agonists) of the activity of YY1, of LSF, of HDAC1
15 or of the transcription repressor complexes containing HDAC1, YY1 and LSF, *e.g.*, but not limited to, antibodies against HDAC1, YY1, LSF or the TRC containing YY1 and LSF; HDAC1, YY1 and/or LSF antisense nucleic acids, organic and inorganic small molecules such as peptides, peptidomimetics, polyamides (*e.g.*, those described by Gottesfeld et al., 1997, *Nature* 387: 202-205), etc. Examples of therapeutics are
20 those proteins, derivatives and analogs of YY1, LSF and HDAC1, as well and inhibitors of the TRC activity, described above. Additional examples are set forth in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

25 A preferred embodiment of the invention relates to methods of using a therapeutic for treatment of HIV infection, preferably HIV-1 infection, in a human subject. The therapeutic of the invention can be used to prevent progression of HIV-1 infection to ARC or to AIDS in a human patient, or to treat a human patient with ARC or AIDS.

30 In a specific embodiment, the therapeutic method of the invention is carried out as monotherapy, *i.e.*, as the only agent provided for treatment of HIV. In another embodiment, the therapeutic is administered in combination with one or more anti-viral compounds, for example, protease inhibitors (*e.g.*, saquinavir, indinavir, ritonavir,

nelfinavir) and/or reverse transcriptase inhibitors (e.g., azidothymidine (AZT), lamivudine (3TC), dideoxyinosine (ddI), dideoxycytidine (ddC), nevirapine, and efavirenz). The therapeutic may also be administered in conjunction with chemotherapy (e.g., treatment with adriamycin, bleomycin, vincristine, vinblastine, doxorubicin and/or Taxol) or other therapies known in the art.

In another embodiment, HIV infection is treated by administration of a therapeutic of the invention in combination with one or more chemokines. In particular, the therapeutic is administered with one or more C-C type chemokines, especially one or more from the group RANTES, MIP-1 α , MIP-1 β and MDC, or the C-X-C type chemokine, SDF-1.

In another embodiment, HIV infection is treated by administration of a combination of one or more transcription factor therapeutics and one or more HIV protein therapeutics of the invention.

In another embodiment, HIV infection is treated by administration of a therapeutic of the invention to antagonize transcriptional repression of the HIV1 LTR. Examples of such therapeutics are discussed parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

One aspect of the invention relates to assaying preparations of YY1 and LSF, and/or derivatives and/or analogs thereof, for efficacy in treatment or prevention of HIV infection. The therapeutic effectiveness of these preparations can be tested by the *in vitro* or *in vivo* assays described in or by any method known in the art for assaying HIV infection, transcription or replication. It is preferable to test the preparation in an *in vitro* assay, e.g., for HIV infection, replication, transcription from the HIV LTR or binding to the HIV LTR by an EMSA, or *in vivo* in an animal model, such as HIV transgenic mice or SIV infected monkeys, before assaying the preparation in humans.

In a specific embodiment, a preparation comprising YY1, LSF and HDAC1 is used.

The HDAC1, YY1 and LSF -related polypeptides are preferably prepared by any chemical or enzymatic synthesis method known in the art, as described above.

The invention provides methods of treatment and prevention by administration to a subject in need of such treatment of a therapeutically or prophylactically effective

amount of a therapeutic of the invention. The subject is preferably an animal, including, but not limited to, animals such as monkeys, cows, pigs, horses, chickens, cats, dogs, etc., and is preferably a mammal, and most preferably human.

Various delivery systems are known and can be used to administer a
5 therapeutic of the invention, *e.g.*, encapsulation in liposomes, microparticles, microcapsules, recombinant cells capable of expressing the therapeutic, receptor-mediated endocytosis (see, *e.g.*, Wu and Wu, 1987, J. Biol. Chem. 262:4429-4432), construction of a therapeutic nucleic acid as part of a retroviral or other vector, etc. Methods of introduction include but are not limited to intradermal, intramuscular,
10 intraperitoneal, intravenous, subcutaneous, intranasal, epidural, and oral routes. The compounds may be administered by any convenient route, for example by infusion or bolus injection, by absorption through epithelial or mucocutaneous linings (*e.g.*, oral mucosa, rectal and intestinal mucosa, etc.) and may be administered together with other biologically active agents. Administration can be systemic or local. In addition,
15 it may be desirable to introduce the pharmaceutical compositions of the invention into the central nervous system by any suitable route, including intraventricular and intrathecal injection; intraventricular injection may be facilitated by an intraventricular catheter, for example, attached to a reservoir, such as an Ommaya reservoir. Pulmonary administration can also be employed, *e.g.*, by use of an inhaler or
20 nebulizer, and formulation with an aerosolizing agent.

Such delivery systems, administration routes, and components thereof are known in the art and described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

25 Where the therapeutic is a nucleic acid encoding a protein therapeutic, the nucleic acid can be administered by gene therapy methods as herein, known in the art and/or described in parent patent application, U.S. Serial No. 09/355,010 and International Publication No. WO 98/33067, mentioned above and incorporated by reference in their entirety herein.

30

Pharmaceutical Compositions:

The present invention also provides pharmaceutical compositions. Such compositions comprise a therapeutically effective amount of a therapeutic, and a pharmaceutically acceptable carrier.

5 The term "pharmaceutically acceptable" means approved by a regulatory agency of the Federal or a state government or listed in the U.S. Pharmacopeia or other generally recognized pharmacopeia for use in animals, and more particularly in humans.

10 The term "carrier" refers to a diluent, adjuvant, excipient, or vehicle with which the therapeutic is administered. Such pharmaceutical carriers can be sterile liquids, such as water and oils, including those of petroleum, animal, vegetable or synthetic origin, such as peanut oil, soybean oil, mineral oil, sesame oil and the like. Water is a preferred carrier when the pharmaceutical composition is administered intravenously. Saline solutions and aqueous dextrose and glycerol solutions can also be employed as
15 liquid carriers, particularly for injectable solutions. Suitable pharmaceutical excipients include starch, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, sodium stearate, glycerol monostearate, talc, sodium chloride, dried skim milk, glycerol, propylene glycol, water, ethanol and the like. The composition, if desired, can also contain minor amounts of wetting or emulsifying agents, or pH buffering
20 agents.

 These compositions can take the form of solutions, suspensions, emulsion, tablets, pills, capsules, powders, sustained-release formulations and the like. The composition can be formulated as a suppository, with traditional binders and carriers such as triglycerides. Oral formulation can include standard carriers such as
25 pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate, etc. Examples of suitable pharmaceutical carriers are described in "Remington's Pharmaceutical Sciences" by E.W. Martin. Such compositions will contain a therapeutically effective amount of the therapeutic, preferably in purified form, together with a suitable amount of carrier so as to provide
30 the form for proper administration to the patient. The formulation should suit the mode of administration.

The therapeutics of the invention can be formulated as neutral or salt forms. Pharmaceutically acceptable salts include those formed with free amino groups such as those derived from hydrochloric, phosphoric, acetic, oxalic, tartaric acids, etc., and those formed with free carboxyl groups such as those derived from sodium, potassium, ammonium, calcium, ferric hydroxides, isopropylamine, triethylamine, 2-ethylamino ethanol, histidine, procaine, etc.

The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical

compositions of the invention. Optionally associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration.

5 In one embodiment, the pharmaceutical compositions of the invention may contain an effective amount of an HDAC1, LSF and/or YY1 antisense nucleic acid in a pharmaceutically acceptable carrier can be administered to a patient having a disease or disorder which is of a type that expresses the YY1-LSF-HDAC1 complexes or HDAC1, LSF or YY1 RNA or protein.

10 The amount of HDAC1, LSF and/or YY1 antisense nucleic acid that will be effective in the treatment of a particular disorder or condition will depend on the nature of the disorder or condition, and can be determined by standard clinical techniques. Where possible, it is desirable to determine the antisense cytotoxicity in vitro, and then in useful animal model systems prior to testing and use in humans.

15 In a specific embodiment, antisense pharmaceutical compositions can be administered via liposomes, microparticles, or microcapsules. In various embodiments of the invention, it may be useful to use such compositions to achieve sustained release of the antisense nucleic acids. In a specific embodiment, it may be desirable to utilize liposomes targeted via antibodies to specific identifiable cell types (Leoneti et al.,
20 1990, *Proc. Natl. Acad. Sci. U.S.A.* 87: 2448-2451; Renneisen et al., 1990, *J. Biol. Chem.* 265: 16337-16342), e.g. to HIV infected T cells.

EXAMPLES

Materials And Methods

25 Nuclear extracts. Large-scale preparation of nuclear extract from CEM cells for chromatographic purification of the TRC were prepared as described (12), with the following minor modifications: buffer A and C were supplemented with 1mM NaF, 1mM Na₂VO₄, 10 µg of leupeptin per ml, 10 µg of aprotinin per ml, 1µg of pepstatin A per ml. Chymostatin (1µg/ml) was also added to buffer A and 50mM beta-glycerophosphate to buffer C.
30

Ion-exchange chromatography. Activated P11 phosphocellulose (Whatman, Clifton, NJ) was equilibrated with 50mM NaCl, 50mM HEPES (pH 7.9), 10%

glycerol, 0.2mM EDTA, 0.5mM phenylmethylsulfonyl fluoride (PMSF) and 0.5mM dithiothreitol (DTT). CEM cell nuclear extract was loaded at 0.4 ml/min, washed, and eluted in a linear gradient of 50mM to 1 M NaCl. Fractions shown by Western blot with anti-YY1 C-20 antibody (Santa Cruz Biotechnology, Santa Cruz, CA) to contain YY1, and shown by EMSA to contain RCS binding activity were pooled and dialyzed against 20mM Tris-HCl (pH 7.9), 10% glycerol, 0.2mM EDTA, 0.5mM PMSF, 0.5mM DTT, and 50mM NaCl before DEAE-cellulose chromatography. A DEAE-cellulose DE52 column (Whatman, Clifton, NJ) was loaded with pooled fractions at 0.2 ml/min. The column was washed and eluted, and fractions analyzed as above. Fractions positive both in Western blot and gel shift were subjected to further purification by DNA-affinity chromatography.

DNA-affinity chromatography. A double stranded oligonucleotide spanning the region -10 to +27 of the HIV-1 LTR was ligated and coupled to CNBr-activated Sepharose CL-4B (Pharmacia, Piscataway, NJ) as previously described (27). Active fractions from DEAE-cellulose chromatography were equilibrated in buffer Z (25mM HEPES [pH 7.6], 0.1 M NaCl, 20% glycerol, 12.5mM MgCl₂, 1mM DTT, 0.5mM PMSF, 0.1% Nonidet P-40). Affinity resin was washed extensively with buffer Z without glycerol and Nonidet P-40. Fractions were incubated for 10 minutes at 4°C with 10 µg of (dI-dC) per ml, loaded by gravity, washed, and eluted with a step gradient of 0.1 to 1 M NaCl. Western blot analysis for detection of histone deacetylase was performed using a rabbit polyclonal antibody raised against a peptide corresponding to the 319-334 C-terminal amino acids of the molecule (59), and against LSF using rabbit polyclonal anti-CP2 antiserum (LSF, LBP-1c; gift of M. Sheffery). Histone deacetylase assays were performed as previously described (31).

Cell lines, transfections and assays. Transfections of HeLa cells were performed as previously described (49). HeLa-CD4-LTR (9) cells were grown in DMEM supplemented with 10% FCS and transfected with 20 µg plasmid DNA (prepared using EndoFree plasmid kit, Qiagen, Valencia, CA) by calcium phosphate coprecipitation as per manufacturer's instructions (ProFection system, Promega, Madison, WI). After 30 minutes at room temperature, the solution was added to the cells (2.5–4x10⁵ cells/plate). Twelve hours after transfection, the cells were washed

with PBS and fed fresh medium. Forty-eight hours later the cells were harvested, cellular extracts prepared, and CAT assays performed as previously described (49). To control for the effect of transcription factor overexpression on general cellular promoters a reporter construct driven by the beta-actin promoter, pH \square -actin-luciferase (66), was used in cotransfection and CAT expression normalized for luciferase activity. Other plasmids used have been previously described (2, 49). Luciferase assays were performed at 48 hours as suggested by the manufacturer (Luciferase assay system, Promega, Madison, WI) but cells resuspended in 200 μ l of lysis buffer and one freeze/thaw step performed. Up to 30 μ l of cellular extract (normalized for protein concentration) in a final volume of 130 μ l was used for luciferase reactions.

For virus production experiments 2×10^4 HeLa cells were transfected with 10 μ l of Superfect (Qiagen, Valencia, CA) and 2.5-3.0 μ g of DNA in a volume of 0.6 ml for 3 hours, washed with PBS, and then grown in 2 ml. Aliquots of culture medium were sampled for detection of HIV-1 p24^{gag} protein by antigen capture enzyme-linked immunosorbent assay according to the manufacturer's instructions (Coulter Corporation, Hialeah, FL).

Immunoprecipitation and electrophoretic mobility shift assays (EMSA):
Immunoprecipitation was performed using nuclear extracts prepared from a Jurkat T cell line (49). 20 μ l samples of extract were mixed with antibody (rabbit IgG, -YY1 {C20; Upstate Biotechnology, Lake Placid, NY} or α -LSF {a gift of M. Sheffery and S. Swendenmann}) at 4°C for 1 hour. 5 μ l α -rabbit IgG agarose conjugated antibody (Sigma, St. Louis, MO) was added and incubation was continued for 1 hour. The complex was precipitated by centrifugation at 3000 rpm, 4°C for 5 minutes. The pellet was washed 3 times with PBS, resuspended in 30 μ l SDS-PAGE sample buffer before separation by SDS-PAGE and Western analysis with α -YY1 or α -LSF antibodies. Bands were visualized using HRP conjugated α -rabbit IgG (Sigma).

Histidine-tagged YY1 (his-YY1) and histidine-tagged LSF (his-LSF) were expressed and harvested in *E. coli* as previously described (53, 64). The RCS oligonucleotide (-10 to +27 of HIV-1 LTR, ref. 49) was end labeled and 1×10^5 cpm was incubated with varying amounts of his-LSF for 20 min at 25°C. GST-YY1 or his-YY1 was added, total protein content normalized by the addition of BSA, and the

reaction continued for 30 minutes. EMSA was then performed as previously described (49). Supershifts were performed by addition of either concentrated α -YY1 (C20-X; Santa Cruz Biotech, Santa Cruz, CA) or α -LSF (64) to the reaction mixture.

In vitro protein interaction mapping: Glutathione-S-transferase (GST)-YY1/GFI-1 chimeras (17) and GST-LSF deletion (54, 55) constructs were expressed and harvested in *E. coli* as previously described. Proteins were visualized by Coomassie brilliant blue staining and protein content was normalized by densitometric analysis. LSF was transcribed and translated *in vitro* by T7 RNA polymerase in rabbit reticulocyte (Promega, Madison, WI) in the presence of 35 S-methionine as per manufacturer's instruction (54). Following capture of GST-YY1/GFI-1 chimera proteins on glutathione-agarose beads, equal volumes of beads were incubated with 35 S-methionine labeled LSF in incubation buffer (17) for 1 hour at 4°C. The beads were washed in incubation buffer/100mM KCl and resuspended in 20 μ l 2x SDS loading buffer. Retained LSF was separated in a 10% SDS gel, dried, and visualized by autoradiography. Similarly, YY1 was transcribed and translated *in vitro*, incubated with captured GST-LSF constructs, and retained YY1 was by autoradiography following electrophoresis.

Example 1:

YY1 and LSF interact *in vivo*

Complex formation at the RCS site can be ablated by either α -YY1 or α -LSF, suggesting that both YY1 and LSF are necessary to form this regulatory complex (49). In this example, we describe evidence that these factors interacted directly in the absence of a DNA binding sequence.

Jurkat CD4⁺ T cell nuclear extracts were incubated with α -YY1, α -LSF, or a nonspecific rabbit polyclonal antiserum. Antibody-protein complexes were precipitated by addition of α -IgG-agarose beads and centrifugation. Precipitates were then assayed for the presence of LSF by Western blot analysis. Immunoprecipitation was specific, as only trace amounts of LSF were recovered by the nonspecific antiserum. α -YY1 precipitated approximately 75% of the LSF activity that could be recovered by α -LSF (Fig. 1a). This indicates that LSF interacts with YY1 *in vivo* in

the absence of the HIV RCS binding site and suggests that direct protein-protein interaction between YY1 and LSF is necessary for complex formation at the HIV LTR.

5 Example 2: LSF and YY1 associate without cofactors *in vitro*

Romerio et al (49) showed that both LSF and YY1 form a complex at the RCS site of the HIV LTR. It was not known whether LSF and YY1 were sufficient to form this complex. We reconstituted this complex *in vitro*, in the absence of other factors to address this question.

10 Histidine-tagged recombinant LSF and YY1 proteins were expressed, harvested from *E. coli*, and allowed to interact with an oligonucleotide encoding the HIV LTR RCS (SEQ ID NO. 1). Up to 100 ng of His-YY1 alone did not form a detectable complex with the RCS (Fig. 1b). Whereas 10 ng of LSF formed a diffuse faintly visible EMSA band (Figs 1b and 1c), the addition of increasing amounts (20
15 ng to 100 ng) of YY1 resulted in increasing levels of complex formation in the presence of 10 ng of LSF (Fig. 1b). As little as 20 ng of His-LSF induced a very prominent protein-DNA complex (Fig. 1c), similar to previous studies (18, 26). Under these conditions, no additional effect of YY1 was discernable when added to 20 ng of LSF. However YY1 specifically enhanced TRC formation, as all reactions were
20 normalized for total protein content by the addition of BSA.

These protein-DNA complexes were supershifted by either α -YY1 or α -LSF antibodies (Fig. 1c), confirming that these complexes contained both LSF and YY1. α -LSF completely shifted the TRC, whereas under these conditions only a fraction of complexes were supershifted by the addition of excess α -YY1. The quantity of α -YY1
25 (5 μ l) completely supershifted YY1 complexes formed on a canonical AAV P5 binding site (data not shown). The effect of the antibody was specific, as α -YY1 had no effect on the mobility of the complex in the absence of YY1 protein (data not shown). Although other factors may be present in the TRC *in vivo*, YY1 and LSF are sufficient to form a complex at the RCS.

30 The addition of YY1 to LSF bound to the RCS was not associated with a further change in the mobility of the DNA-protein complex.

Example 3: Interaction domains of YY1 and LSF

The interaction of LSF with YY1 was mapped using constructs that contained a series of nested deletions within the coding region of LSF (**Fig. 2a**).

Neither the carboxyl nor amino terminus of LSF was required for interaction
5 with YY1. However, the central region of the protein was required for interaction with YY1, as amino terminal deletions beyond amino acid 164, and carboxy terminal deletions prior to amino acid 403 resulted in marked diminution of the ability of LSF to bind YY1 (**Fig. 2b**). Binding was lost altogether when c-terminal sequences between amino acid residues 308 and 368 were removed. Further amino acid
10 substitutions within this region, which impair multimerization (55), markedly decreased the ability of the mutant LSF to bind full length YY1. A likely possibility is that LSF recognizes YY1 only in its multimeric conformation.

YY1 DNA-binding activity and many YY1-protein interactions map to the carboxyl-terminal zinc fingers of the molecule. Therefore, YY1 interaction domains
15 were mapped using chimeric YY1 recombinants (**Fig. 2c**). These chimeras expressed GST fused to the N-terminal domain of the protein and had varying numbers of YY1 zinc finger domains replaced by the structurally homologous GFI-1 zinc fingers (17). No deleterious effect on function or stability of YY1 was observed (17). **Figure 2c** shows that the interaction with LSF the third and fourth zinc fingers of YY1 are not
20 required to retain LSF. LSF bound *in vitro* to constructs that contained YY1 zinc fingers 1, 2, or both. In these assays chimera 2, which expresses zinc fingers 1 and 2 bound LSF more avidly than intact YY1. Chimera 1 bound LSF nearly as well as YY1, while chimera 7 could bind LSF *in vitro*, but less avidly. Thus, either zinc finger allowed binding to LSF, but binding was optimal when both fingers were present.
25 Chimera 5, containing only the last two zinc fingers of YY1, did not bind LSF. The GST-Y/GFI-1 construct containing the entire YY1 amino-terminal domain fused to the zinc finger domain of the GFI-1 protein retained minimal amounts of LSF. Finally, a chimera expressing only the YY1 zinc finger domains, and lacking the entire amino terminal YY1 region, bound LSF at least as well as intact YY1. Therefore, YY1
30 requires only zinc fingers 1 and 2 to recognize LSF.

Example 4:

LSF competent to bind DNA is required for repression of HIV LTR expression

Shirra and Hansen (54) and Shirra et al (55) demonstrated that LSF binds a canonical SV40 late promoter site via the formation of homotetramers. Further, binding can be blocked by the expression of a dominant negative mutant defective in DNA binding but with remaining ability to multimerize (LSF 234QL/236KE or dnLSF). The following experiments were performed to test the effect of dnLSF using the HeLa-CD4-LTR cell line (9).

The LTR reporter carried by HeLa-CD4-LTR cell line exists within the native chromatin structure of the genome. Transfection of YY1 inhibited Tat-activated CAT activity in these cells (**Fig. 3**) in agreement with previous studies using plasmid-based reporters (49). In the setting of a chromosomally integrated reporter gene the provision of LSF augmented repression mediated by YY1, confirming the effect of YY1 and LSF on an integrated HIV-1 promoter. Significantly, dnLSF abolished the ability of YY1 to repress CAT expression, confirming that LSF capable of binding DNA is required to allow YY1 to repress HIV-1 LTR expression (**Fig. 3**). As in previous studies (49), these effects were specific to the HIV LTR; results are normalized to the expression of a cotransfected beta-actin-luciferase reporter gene, whose expression was not significantly affected by YY1, LSF, or dnLSF (not shown).

Example 5: Copurification of HDAC1 with the YY1-LSF complex

YY1 has been demonstrated to act via the recruitment of histone deacetylases (71, 72). As a nucleosome is present near the RCS when the HIV-1 LTR is integrated (46, 51, 61), we tested to see if histone deacetylase was present in the RCS DNA affinity chromatography fractions.

The YY1-LSF complex was purified by phosphocellulose P11, DEAE-cellulose column, and DNA affinity chromatography, as previously described (49). YY1, LSF and the RCS-binding activity copurified in the 0.3 and 0.4 M NaCl fractions of the final step of purification (**Fig. 4a**). RCS-binding activity was enriched about 10,000-fold by this procedure. As shown in **Fig. 4a**, a rabbit polyclonal antibody raised against amino acids 319-334 of HDAC1 was able to detect a protein with apparent molecular mass of 66 kDa in the 0.3 and 0.4 M NaCl pooled fractions.

HDAC1, a 55 kDa protein, migrates at this apparent molecular mass in our gel system (58b).

To rule out the possibility that these fractions contained a protein immunologically similar but enzymatically unrelated to histone deacetylase, we assayed the histone deacetylase activity of the DNA affinity chromatography fractions. As expected, the 0.3 and 0.4 NaCl fractions showed strong HDAC1 activity, as measured by release of ^3H -acetic acid (Fig. 4b). These results indicate that active HDAC1 copurifies with the YY1-LSF complex, and suggests that the YY1-LSF complex represses HIV-1 transcription via the recruitment of HDAC1.

Example 6:

Repression of the HIV-1 LTR by YY1 requires interaction with a histone deacetylase

A recent report has demonstrated that the Gly/Ala-rich domain of YY1 mediates the interaction with the histone deacetylase, and is required for repression of the adeno-associated virus (AAV) P5 promoter by YY1 (71). To test whether the mechanism of repression of the HIV-1 LTR by YY1 is mediated by a histone deacetylase, we performed a series of transient transfection experiments using a mutant YY1 deleted of the Gly/Ala-rich domain (YY1 Δ 154-199; ref. 2) required for interaction with HDAC1.

Initial experiments were performed as previously (41, 49), demonstrating that cotransfection of YY1 inhibited Tat-activated, LTR-driven CAT expression. However, YY1 Δ 154-199 was unable to inhibit CAT activity, indicating the absolute requirement of the Gly/Ala-rich domain of YY1 for efficient repression of the HIV-1 LTR (data not shown). Indeed, YY1 Δ 154-199 activated LTR expression, suggesting ongoing competition between constitutive cellular YY1 and HIV LTR activating factors.

Chromatin remodeling effects on gene activity have often been imputed in studies using transfected, plasmid-encoded reporter genes that may not reflect the activity of genes contained in native chromatin. However the LTR reporter carried by the HeLa-CD4-LTR cell line (9) exists within the native chromatin structure of the genome. Significantly, YY1 Δ 154-199 failed to repress CAT expression, confirming that the Gly/Ala-rich histone deacetylase interaction domain is required for repression

of the HIV-1 LTR by YY1 (Fig. 3). Indeed, YY1 Δ 154-199 activated CAT expression, but when normalized for modest activation of a cotransfected beta-actin-luciferase reporter, this effect was not significant. Repression was also blocked by addition of the specific HDAC1 inhibitor trichostatin-A to the culture medium (data not shown).

This is the first demonstration of both cooperative repression by YY1 and LSF, and the lack of repression by YY1 Δ 154-199 in the context of a chromosomally integrated HIV-1 promoter. It is also the first demonstration in this context that YY1 requires its HDAC1 interaction domain to mediate repression.

Example 7: Repression of the HIV-1 virion production by YY1 requires interaction with a histone deacetylase

Support for the role of HDAC1 in repression of HIV-1 virion production was demonstrated by cotransfection of HeLa cells with the infectious molecular clones pNL4-3 (1) or pYU-2 (37) and empty CMV vector, CMV-YY1, or CMV-YY1 Δ 154-199. As these cells support HIV replication but cannot be infected, a measurement of the effect of YY1 on a single round of viral replication can be made.

The influence of YY1 on viral production was assayed by testing of culture supernatant for the presence of the viral protein p24^{gag}. Cotransfection with a vector expressing YY1 produced dose-dependent inhibition of either CXCR4 (pNL4-3) or CCR5 (pYU-2) tropic virus, whereas cotransfection of YY1 Δ 154-199 failed to inhibit HIV production (Fig. 5). Again, YY1 Δ 154-199 activated HIV expression above normal levels. Similar results were seen in the CD4⁺ T cell line CEM when transfected with pNL4-3 or pYU-2 and empty CMV vector, CMV-YY1, or CMV-YY1 Δ 154-199 (data not shown). These findings suggest the possibility of ongoing competition between constitutive cellular YY1 and HIV LTR activating factors, although secondary activating effects of CMV-YY1 Δ 154-199 cannot be excluded.

While the invention has been described in conjunction with examples thereof, it is to be understood that the foregoing description is exemplary and explanatory in nature, and is intended to illustrate the invention and its preferred embodiments. Through routine experimentation, the artisan will recognize apparent modifications

and variations that may be made without departing from the spirit of the invention. Thus, the invention is intended to be defined not by the above description, but by the following claims and their equivalents.

All references cited herein are incorporated by reference in their entirety.

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WHAT IS CLAIMED IS:

1. A method of repressing HIV transcription in a human subject in need of such treatment comprising administering to the subject an effective amount of a preparation of a transcription repressor complex (TRC) comprising YY1 or a derivative or analog thereof; LSF or a derivative or analog thereof; and HDAC1 or a derivative or analog thereof.
2. A method of repressing HIV transcription in a human subject in need of such treatment comprising administering to the subject an effective amount of an agent that affects the activity of a transcription repressor complex (TRC), said complex comprising YY1 or a derivative or analog thereof; LSF or a derivative or analog thereof; and HDAC1 or a derivative or analog thereof.
3. The method of claim 2 wherein said agent enhances the binding of a YY1 or a derivative or analog thereof to HDAC1 or a derivative or analog thereof.
4. The method of claim 2 wherein said agent enhances recruitment of HDAC1 or a derivative or analog thereof by YY1 or a derivative or analog thereof.
5. The method of claim 2 wherein said agent enhances the enzymatic activity of HDAC1 or a derivative or analog thereof.
6. The method of claim 2 wherein said agent enhances the expression of HDAC1 or a derivative or analog thereof.
7. The method of claim 2 wherein said agent up-regulates the expression of HDAC1 or a derivative or analog thereof.
8. The method of claim 2 wherein said agent comprises an effective amount of a nucleic acid or combination of nucleic acids comprising nucleotide sequences encoding YY1 or a derivative or analog thereof; nucleotide sequences encoding LSF or a derivative or analog thereof; and nucleotide sequences encoding HDAC1 or a derivative or analog thereof.
9. A method of treating quiescent reservoirs of HIV infection in a human subject in need of such treatment comprising:
 - (a) administering to the subject an amount of an agent that down-regulates the expression of HDAC1 or a derivative or analog thereof, the amount effective to down-regulate transcription repressor complex-associated repression of HIV transcription;

(b) allowing latent, quiescent reservoirs of HIV to become actively transcribing; and

(c) treating the subject with an effective amount of an antiretroviral agent.

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10. A method of treating quiescent reservoirs of HIV infection in a human subject in need of such treatment comprising:

(a) administering to the subject an amount of an agent that inhibits the expression of HDAC1 or a derivative or analog thereof, the amount effective to inhibit transcription repressor complex-associated repression of HIV transcription;

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(b) allowing latent, quiescent reservoirs of HIV to become actively transcribing; and

(c) treating the subject with an effective amount of an antiretroviral agent.

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11. A method of treating or preventing latent HIV infection in a human subject in need of such treatment or prevention comprising administering to the subject (a) an amount of an inhibitor of an HDAC1-recruiting activity of YY1, said amount being effective to inhibit repression of HIV transcription, and (b) a therapeutically effective amount of one or more anti-viral drugs selected from the group consisting of AZT, 3TC, ddI, ddC, 3TC, saquinavir, indinavir, ritonavir, nelfinavir, nevirapine and efavirenz.

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12. An isolated, purified composition comprising a transcription repressor complex (TRC) comprising YY1; LSF and HDAC1, or derivatives or analogs thereof.

13. A method for identifying agents having HIV transcription repressing activity comprising screening a compounds library using the composition of claim 11 to identify analogs thereof and assaying said analogs for transcription repressing activity.

14. A method for modulating HIV transcription in a human subject in need thereof comprising administering to said subject a compound that modulates the histone structure of the RCS site of the HIV LTR and, therefore, modulates the activity of the transcription repressor complex.

15. A method for modulating HIV transcription in a human subject in need thereof comprising administering to said subject a compound that modulates the association between YY1 and HDAC1 at the RCS site of the HIV LTR and, therefore, modulates the activity of the transcription repressor complex.
16. A pharmaceutical composition comprising an effective amount of a nucleic acid or combination of nucleic acids comprising nucleotide sequences encoding YY1 or a derivative or analog thereof; nucleotide sequences encoding LSF or a derivative or analog thereof; and nucleotide sequences encoding HDAC1 or a derivative or analog thereof, said amount effective to repress HIV transcription, thereby treat HIV infection.
17. A pharmaceutical composition comprising an effective amount of an agent that affects the activity of a transcription repressor complex (TRC), said complex comprising YY1 said complex comprising YY1 or a derivative or analog thereof; LSF or a derivative or analog thereof; and HDAC1 or a derivative or analog thereof and said amount being effective to repress HIV transcription and thereby treat HIV infection.
18. The pharmaceutical composition of claim 17 wherein said agent enhances the binding of a YY1 or a derivative or analog thereof to HDAC1 or a derivative or analog thereof.
19. The pharmaceutical composition of claim 17 wherein said agent enhances recruitment of HDAC1, or a derivative or analog thereof, by YY1 or a derivative or analog thereof.
20. The pharmaceutical composition of claim 17 wherein said agent enhances the enzymatic activity of HDAC1 or a derivative or analog thereof.
21. The pharmaceutical composition of claim 17 wherein said agent enhances the expression of HDAC1 or a derivative or analog thereof.
22. The pharmaceutical composition of claim 17 wherein said agent up-regulates the expression of HDAC1 or a derivative or analog thereof.
23. A pharmaceutical composition comprising an effective amount of YY1, LSF and HDAC1, said amount effective to repress HIV transcription, thereby treat HIV infection.

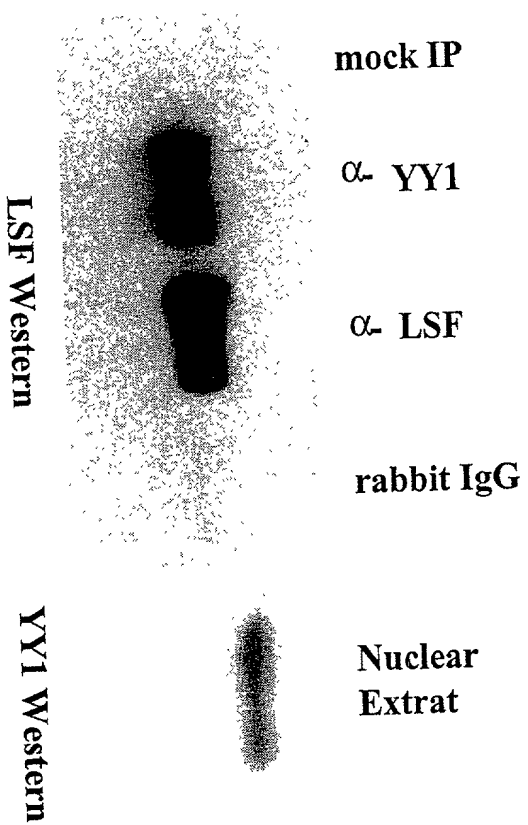
**AN HIV TRANSCRIPTION REPRESSOR COMPLEX AND COMPOSITIONS
AND METHODS BASED THEREON**

ABSTRACT

The molecular mechanism of YY1/LSF-associated repression of HIV-1 is described herein. More particularly, an HIV transcription repressor complex containing YY1, LSF and HDAC1 is described. The invention is based on our discovery that (1) HDAC1 co-purifies with the LTR-binding YY1-LSF repressor
5 complex; (2) the domain of YY1 that interacts with HDAC1 is required to repress the HIV-1 promoter; (3) the expression of HDAC1 augments repression of the LTR by YY1, and (4) the deacetylase inhibitor trichostatin-A blocks repression mediated by YY1. This novel link between HDAC1 recruitment and inhibition of HIV-1 expression
10 by YY1 and LSF, in the natural context of a viral promoter integrated into chromosomal DNA, supports novel HIV therapies described herein and has significant implications for the long-term treatment of AIDS.

-5 Vnd 1/2 page figure Feb 2000

Fig 1a



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Fig 1b

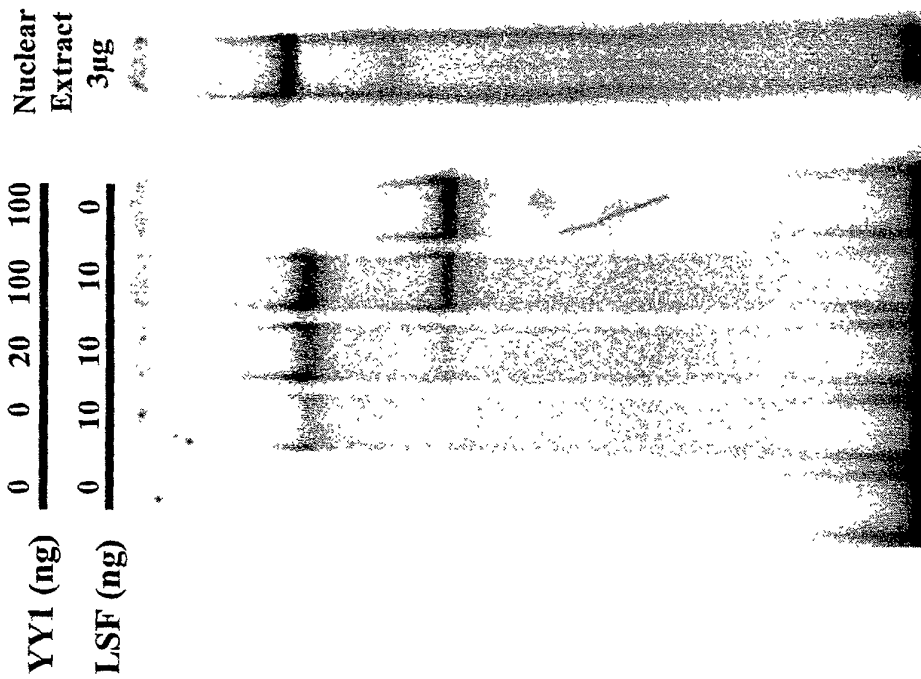


Fig. 1c

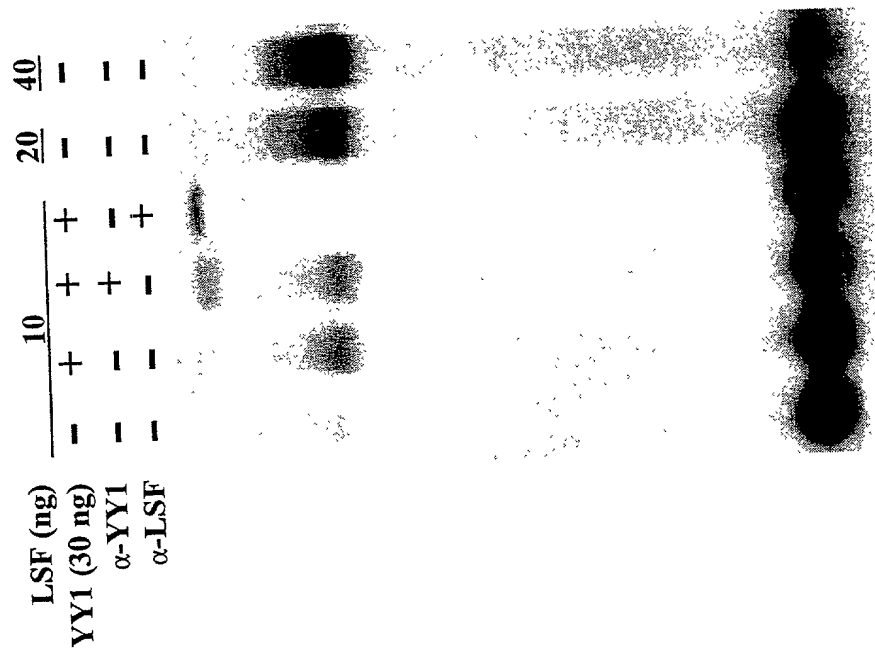


Fig 2a

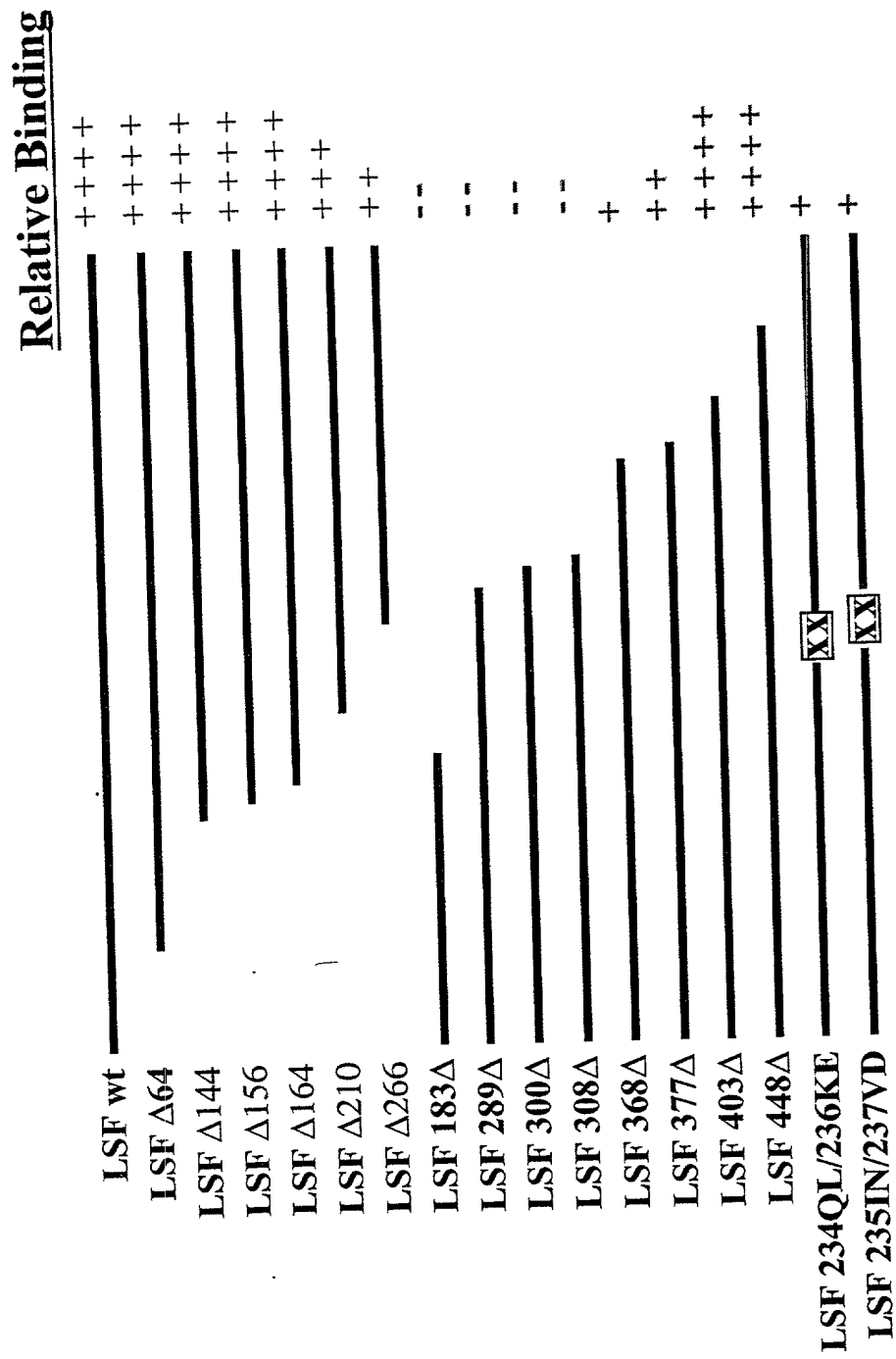


Fig 2b

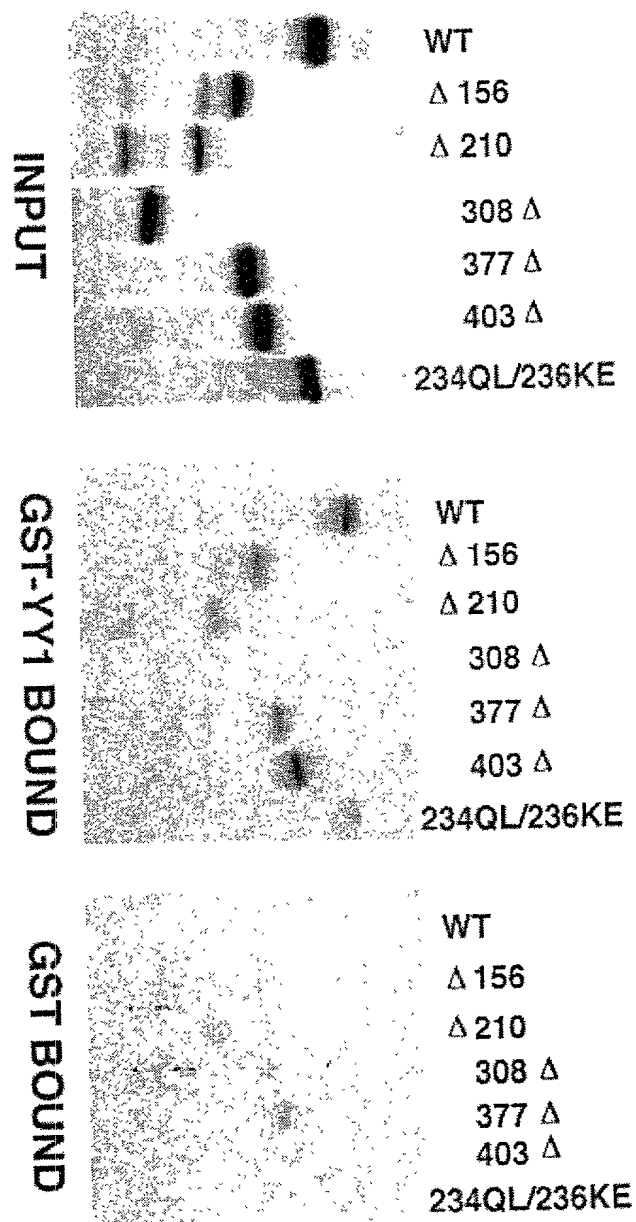


Fig 2c

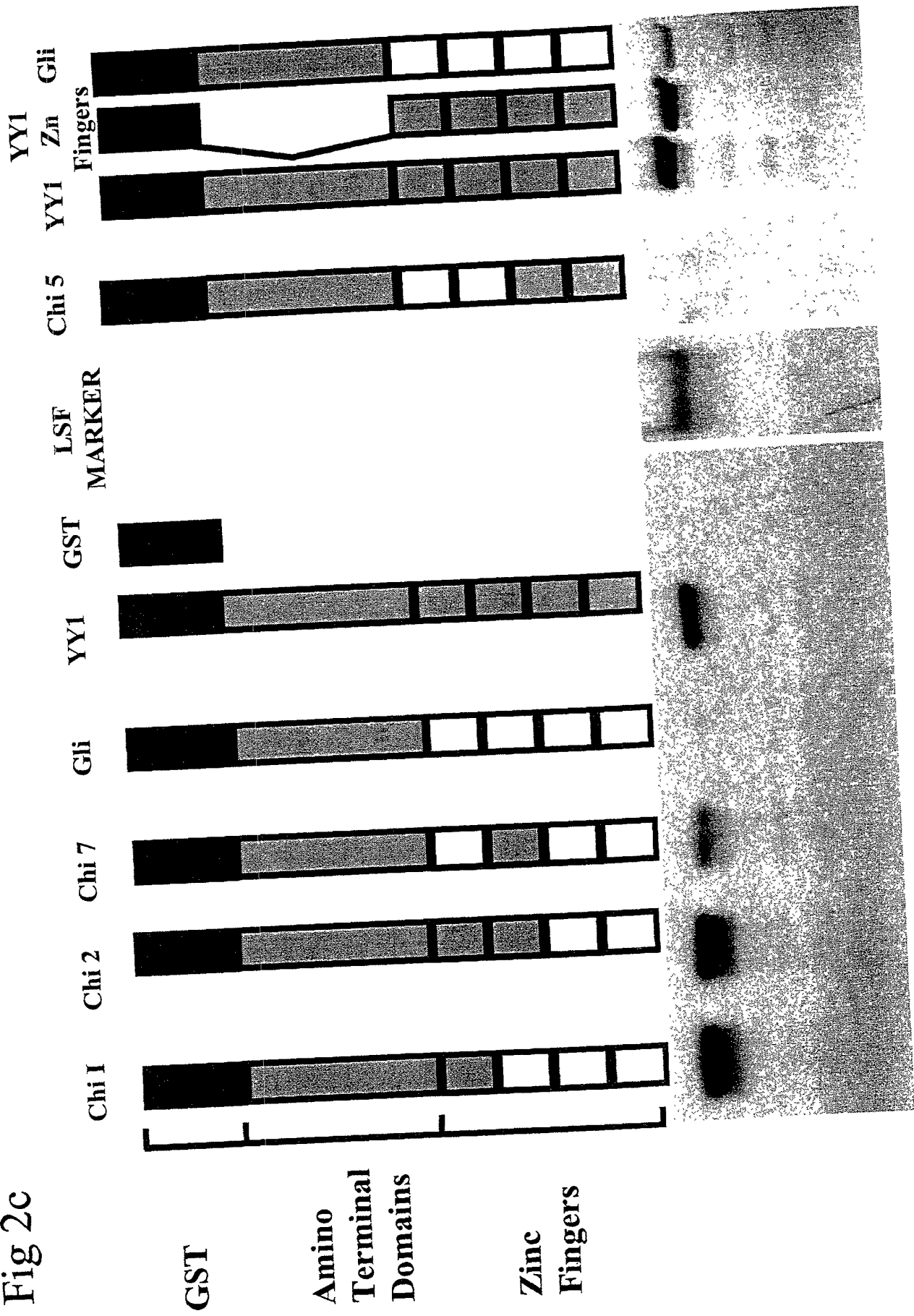


Fig 3

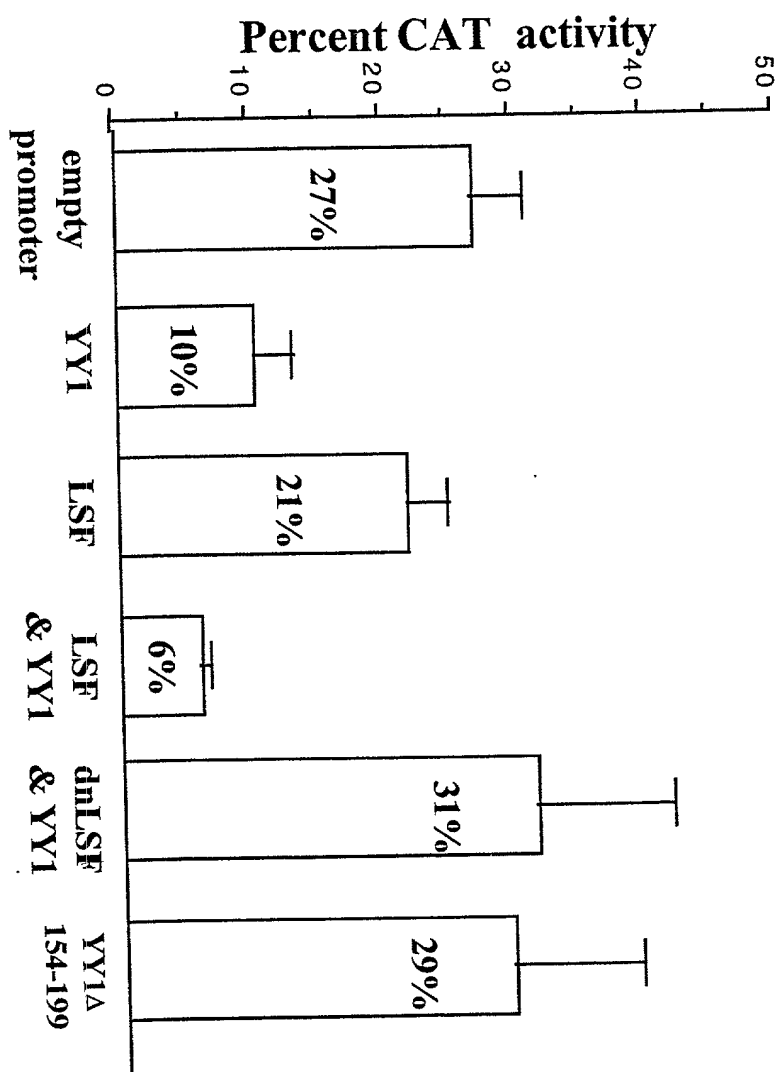


Fig. 4a

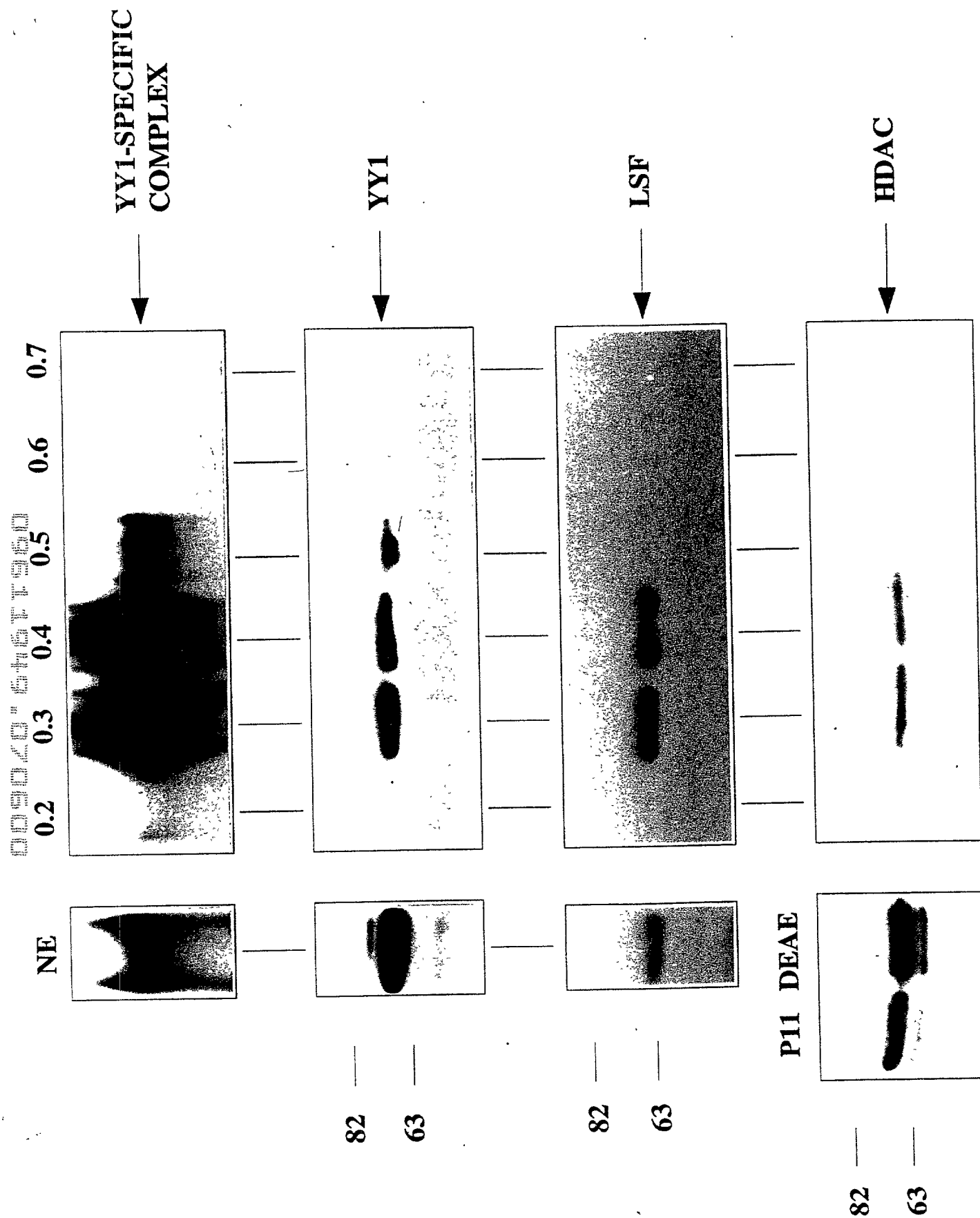


Fig 4b

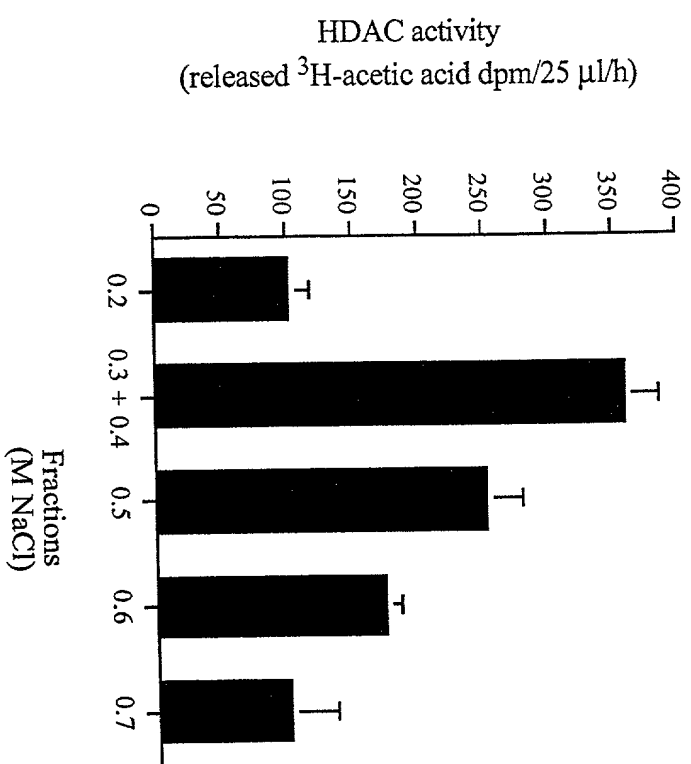


Fig 5

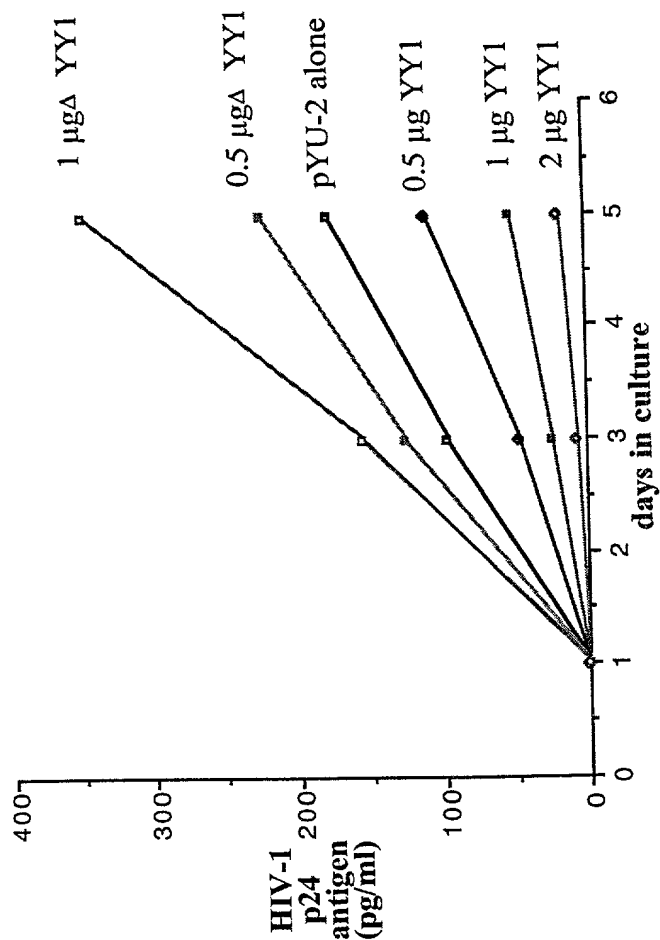
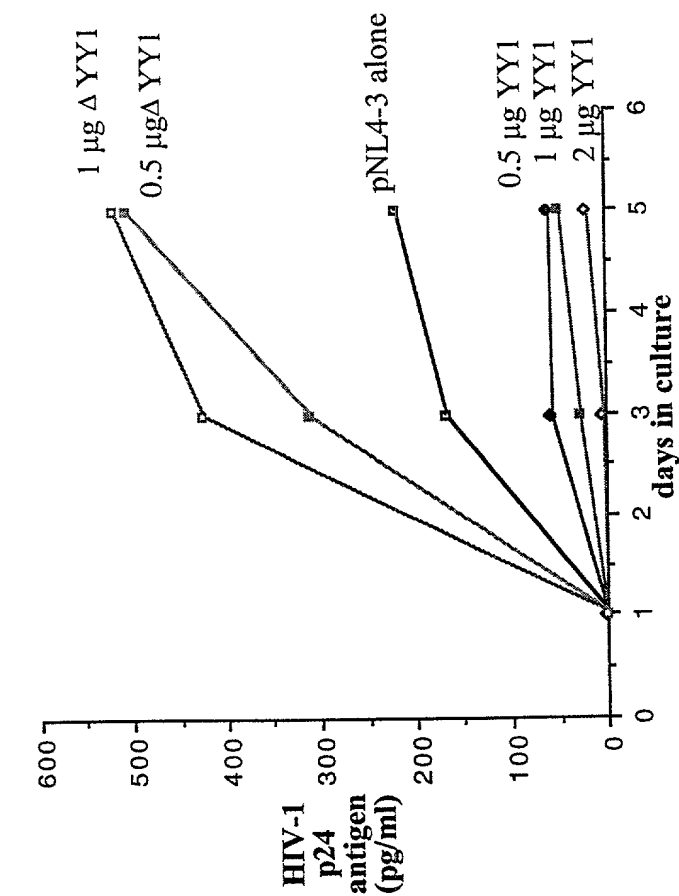
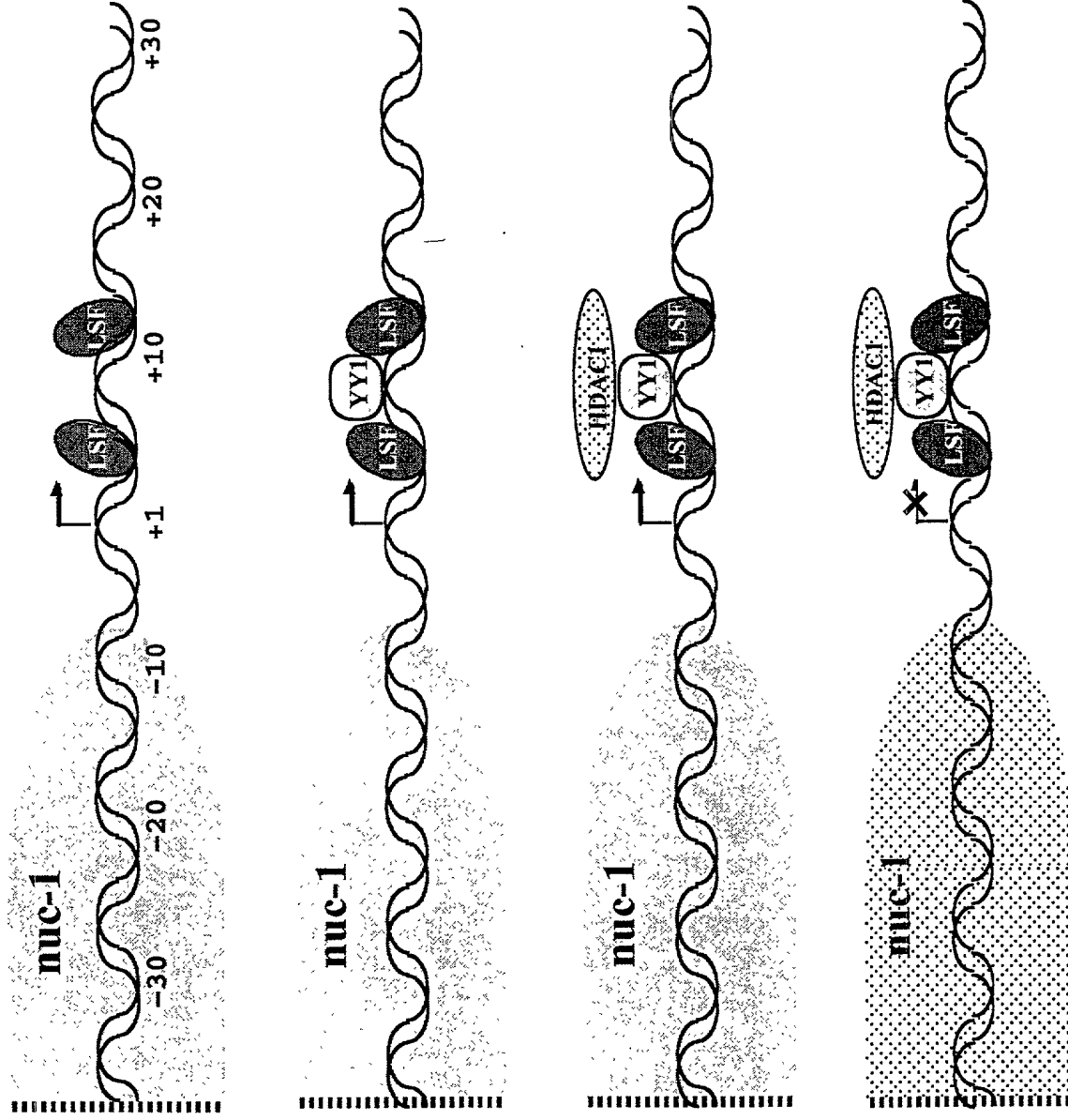


Fig 6

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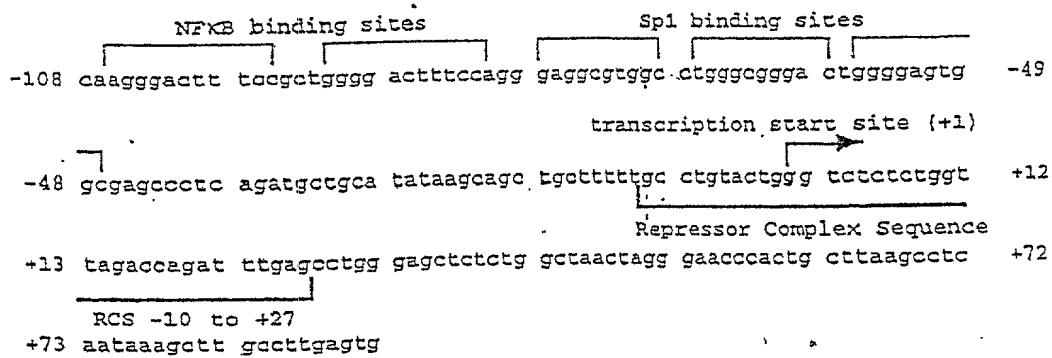


FIG. 7

CGCCGAGACG AGCAGCGGCC GAGCGAGCGC GGGCGCGGGC GCACCGAGGC GAGGGAGGCG	60
GGGAAGCCCC GCGCGCGCGC CCCC GCCCGC CCCTTCCCCC GCGCGCGGCC CCCTCTCCCC	120
CCGCCCCGCTC GCGCGCTTCC TCCCTCTGCC TTCCTTCCCC ACGGCCGGCC GCCTCCTCGC	180
CCGCCCCGCC GCAGCCGAGG AGCCGAGGCC GCGCGCGCGC TGGCGGCGGA GCCCTCAGCC	240
ATG GCC TCG GGC GAC ACC CTC TAC ATC GCC ACG GAC GGC TCG GAG ATG	288
Met Ala Ser Gly Asp Thr Leu Tyr Ile Ala Thr Asp Gly Ser Glu Met	
1 5 10 15	
CCG GCC GAG ATC GTG GAG CTG CAC GAG ATC GAG GTG GAG ACC ATC CCG	336
Pro Ala Glu Ile Val Glu Leu His Glu Ile Glu Val Glu Thr Ile Pro	
20 25 30	
GTG GAG ACC ATC GAG ACC ACA GTG GTG GGC GAG GAG GAG GAG GAG GAC	384
Val Glu Thr Ile Glu Thr Thr Val Val Gly Glu Glu Glu Glu Glu Asp	
35 40 45	
GAC GAC GAC GAG GAC GGC GGC GGT GGC GAC CAC GGC GGC GGC GGC GGC	432
Asp Asp Asp Glu Asp Gly Gly Gly Gly Asp His Gly Gly Gly Gly Gly	
50 55 60	
CAC GGC CAC GCC GGC CAC CAC CAC CAC CAC CAT CAC CAC CAC CAC CAC	480
His Gly His Ala Gly His His His His His His His His His His His	
65 70 75 80	
CCG CCC ATG ATC GCT CTG CAG CCG CTG GTC ACC GAC GAC CCG ACC CAG	528
Pro Pro Met Ile Ala Leu Gln Pro Leu Val Thr Asp Asp Pro Thr Gln	
85 90 95	
GTG CAC CAC CAC CAG GAG GTG ATC CTG GTG CAG ACG CGC GAG GAG GTG	576
Val His His His Gln Glu Val Ile Leu Val Gln Thr Arg Glu Glu Val	
100 105 110	
GTG GGC GGC GAC GAC TCG GAC GGC CTG CGC GCC GAG GAC GGC TTC GAG	624
Val Gly Gly Asp Asp Ser Asp Gly Leu Arg Ala Glu Asp Gly Phe Glu	
115 120 125	
GAT CAG ATT CTC ATC CCG GTG CCC GCG CCG GCC GGC GGC GAC GAC GAC	672
Asp Gln Ile Leu Ile Pro Val Pro Ala Pro Ala Gly Gly Asp Asp Asp	
130 135 140	
TAC ATT GAA CAA ACG CTG GTC ACC GTG GCG GCG GCC GGC AAG AGC GGC	720
Tyr Ile Glu Gln Thr Leu Val Thr Val Ala Ala Ala Gly Lys Ser Gly	
145 150 155 160	
GGC GGC GGC TCG TCG TCG TCG GGA GGC GGC CGC GTC AAG AAG GGC GGC	768
Gly Gly Gly Ser Ser Ser Ser Gly Gly Gly Arg Val Lys Lys Gly Gly	
165 170 175	
GGC AAG AAG AGC GGC AAG AAG AGT TAC CTC AGC GGC GGC GCC GGC GCG	816
Gly Lys Lys Ser Gly Lys Lys Ser Tyr Leu Ser Gly Gly Ala Gly Ala	
180 185 190	
GCG GGC GGC CGC GGC GCC GAC CCG GGC AAC AAG AAG TCG GAG CAG AAG	864
Ala Gly Gly Arg Gly Ala Asp Pro Gly Asn Lys Lys Trp Glu Gln Lys	
195 200 205	
CAG GTG CAG ATC AAG ACC CTG GAG GGC GAG TTC TCG GTC ACC ATG TGG	912
Gln Val Gln Ile Lys Thr Leu Glu Gly Glu Phe Ser Val Thr Met Trp	
210 215 220	
TCC TCA GAT GAA AAA AAA GAT ATT GAC CAT GAG ACA GTG GTT GAA GAA	960
Ser Ser Asp Glu Lys Lys Asp Ile Asp His Glu Thr Val Val Glu Glu	
225 230 235 240	
CAG ATC ATT GGA GAG AAC TCA CCT CCT GAT TAT TCA GAA TAT ATG ACA	1008
Gln Ile Ile Gly Glu Asn Ser Pro Pro Asp Tyr Ser Glu Tyr Met Thr	
245 250 255	

GAGGACGCCA	TGATTGGTGG	GCGCTGGGGC	GGCGGACGGT	GGAAGGGCCT	GCGGAGTCTA	60
GGTTTTACGC	CTGTGCTGGA	CTTTCTCCTT	CCATGTTTCC	AGGCCGTGGG	GGGCTACAGA	120
GGGCGAGAAG	TGCGCTCAGC	GGAAACCTGG	ATTTGGTTCT	AAGCCGTGGG	GTGAGAAGG	180
GGTGACCCGA	AGTGATCGTG	GGAOTGACCG	GAAGCGAGGC	CTGGAGGGGA	AAGAGAGAGC	240
GAGACCTGGG	AGGGAGGGGG	CCTCCAGCAG	AAAGGGGCGG	GGGAAAAGGT	GCAAAAGCAG	300
CGTGGGAGCG	CCGGGCTGGC	TTCCTGCGGC	TGCTGCTGGT	CTGACTGGGA	AGCAGCAAGC	360
CACCACTACG	AACTCTCAAG	AGGAGTGGGA	GTGCGGGAGT	CCAGAGCTGC	CTCTGGGAAG	420
TCTGCAGTAG	TTGAGCAAG	GGGTCTCAC	GTTCTTGAGA	GCTGGGCAGG	GGGGATTG	480
GAACCTGGGG	CAGCCAAGAA	CGAGCAGCCA	AGGGTACGGG	AGATTAGTTG	TGCACAGAGC	540
AGTGCTGGTC	GGGCTTGGGG	GTGGCTGGTG	GGCACTGCGT	GGGAAACCTT	GGTTTGTAGT	600
TTTCTTGGTT	TGCGTTACTC	CTGTTGGGTA	GAATTACCCT	CCGCGCCTTT	GTACAAGACA	660
CGGTGTCTCC	TGGGGCAAGG	AAGGAGCCAG	G	ATG GCC TGG GCT CTG AAG CTG		712
			Met Ala Trp Ala Leu Lys Leu			
			1	5		
CCT CTG GCC GAC GAA GTG ATT GAA TCC GGG TTG GTG CAG GAC TTT GAT						760
Pro Leu Ala Asp Glu Val Ile Glu Ser Gly Leu Val Gln Asp Phe Asp						
	10		15		20	
GCT AGC CTG TCC GGG ATC GGC CAG GAA CTG GGT GCT GGT GCC TAT AGC						808
Ala Ser Leu Ser Gly Ile Gly Gln Glu Leu Gly Ala Gly Ala Tyr Ser						
	25		30		35	
ATG AGT GAT GTC CTT GCA TTG CCC ATT TTT AAG CAA GAA GAG TCG AGT						856
Met Ser Asp Val Leu Ala Leu Pro Ile Phe Lys Gln Glu Glu Ser Ser						
	40		45		50	55
TTG CCT CCT GAT AAT GAG AAT AAA ATC CTG CCT TTT CAA TAT GTG CTT						904
Leu Pro Pro Asp Asn Glu Asn Lys Ile Leu Pro Phe Gln Tyr Val Leu						
	60		65		70	
TGT GCT GCT ACC TCT CCA GCA GTG AAA CTC CAT GAT GAA ACC CTA ACG						952
Cys Ala Ala Thr Ser Pro Ala Val Lys Leu His Asp Glu Thr Leu Thr						
	75		80		85	
TAT CTC AAT CAA GGA CAG TCT TAT GAA ATT CGA ATG CTA GAC AAT AGG						1000
Tyr Leu Asn Gln Gly Gln Ser Tyr Glu Ile Arg Met Leu Asp Asn Arg						
	90		95		100	
AAA CTT GGA GAA CTT CCA GAA ATT AAT GGC AAA TTG GTG AAG AGT ATA						1048
Lys Leu Gly Glu Leu Pro Glu Ile Asn Gly Lys Leu Val Lys Ser Ile						
	105		110		115	
TTC CGT GTG GTG TTC CAT GAC AGA AGG CTT CAG TAC ACT GAG CAT CAG						1096
Phe Arg Val Val Phe His Asp Arg Arg Leu Gln Tyr Thr Glu His Gln						
	120		125		130	135
CAG CTA GAG GGC TGG AGG TGG AAC CGA CCT GGA GAC AGA ATT CTT GAC						1144
Gln Leu Glu Gly Trp Arg Trp Asn Arg Pro Gly Asp Arg Ile Leu Asp						
	140		145		150	
ATA GAT ATC CCG ATG TCT GTG GGT ATA ATC GAT CCT AGG GCT AAT CCA						1192
Ile Asp Ile Pro Met Ser Val Gly Ile Ile Asp Pro Arg Ala Asn Pro						
	155		160		165	
ACT CAA CTA AAT ACA GTG GAG TTC CTG TGG GAC CCT GCA AAG AGG ACA						1240
Thr Gln Leu Asn Thr Val Glu Phe Leu Trp Asp Pro Ala Lys Arg Thr						
	170		175		180	
TCT GTG TTT ATT CAG GTG CAC TGT ATT AGC ACA GAG TTC ACT ATG AGG						1288
Ser Val Phe Ile Gln Val His Cys Ile Ser Thr Glu Phe Thr Met Arg						
	185		190		195	
AAA CAT GGC GGA GAA AAG GGG GTG CCA TTC CGA GTA CAA ATA GAT ACC						1336
Lys His Gly Gly Glu Lys Gly Val Pro Phe Arg Val Gln Ile Asp Thr						
	200		205		210	215

(SEQ ID NO: 6)

[NCBI Gen Bank Accession No. NP 004955]

1 maqtqgtrrk vcyyydgdvg nyyygqghpm kphirmthn lllynglyrk meiyprhkan
61 aeemtkyhsd dyikflrsir pdnmseyskq mqrfnvgedc pvfdglfefe qlstggsvas
121 avklnkqgtd iavnwagglh hakkseasgf cyvndivlai lellkyhgrv lyididihhg
181 dgveeafytt drvmtvsfhk ygeyfgtgd lrdigagkgk yyavnyplrd giddesyeai
241 fkpvmaskvme mfpqsavvlq cgsdslsgdr lgcfnlitkg hakcvefvks fnlpmlmlgg
301 ggytirnvar crtyetaval dteipnelpy ndyfeyfgpd fklhispsnm tnqntneyle
361 kikqrlfenl rmlphapgvg mqaipedaip eesgdededd pdkrisicss dkriaceeef
421 sdseegeggy rknssnfkka krvtkedeke kdpeekkevt eeektkeekp eakgvkeevk
481 la

FIGURE 10B
(SEQ ID NO: 7)
Human Histone Deacetylase 1 mRNA Sequence
[NCBI Gen Bank Accession No. NP 004964]

ORIGIN

1 gagcggagcc gcgggcggga gggcggacgg accgactgac ggtagggacg ggagggcagc
61 aagatggcgc agacgcaggg caccgcgagg aaagtctgtt actactacga cggggatgtt
121 ggaaattact attatggaca aggccacca atgaagcctc accgaatccg catgactcat
181 aatttgctgc tcaactatgg tctctaccga aaaatggaaa tctatcgccc tcacaaagcc
241 aatgctgagg agatgaccaa gtaccacagc gatgactaca ttaaattctt gcgctccatc
301 cgtccagata acatgtcgga gtacagcaag cagatgcaga gattcaacgt tggtgaggac
361 tgtccagtat tcgatggcct gtttgagtgc tgtcagttgt ctactggtgg ttctgtggca
421 agtgctgtga aacttaataa gcagcagacg gacatcgctg tgaattgggc tgggggcctg
481 caccatgcaa agaagtccga ggcctctggc ttctgttacg tcaatgatat cgtcttggcc
541 atcctggaac tgctaaagta tcaccagagg gtgctgtaca ttgacattga tattcaccat
601 ggtgacggcg tggaagaggc cttctacacc acggaccggg tcatgactgt gtcctttcat
661 aagtatggag agtacttccc aggaactggg gacctacggg atatcggggc tggcaaaggc
721 aagtattatg ctgttaacta cccgctccga gacgggattg atgacgagtc ctatgaggcc
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841 cagtgtggct cagactccct atctggggat cggttagggt gcttcaatct aactatcaaa
901 ggacacgcca agtgtgtgga atttgtcaag agctttaacc tgcctatgct gatgctggga
961 ggcgggtggt acaccattcg taacgttgcc cgggtgcagga catatgagac agctgtggcc
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1081 gatttcaagc tccacatcag tccttccaat atgactaacc agaacacgaa tgagtacctg
1141 gagaagatca aacagcgact gtttgagaac cttagaatgc tgccgcacgc acctggggtc
1201 caaatgcagg cgattcctga ggacgccatc cctgaggaga gtggcgatga ggacgaagac
1261 gaccctgaca agcgcactct gatctgctcc tctgacaaac gaattgcctg tgaggaagag
1321 ttctccgatt ctgaagagga gggagagggg ggccgcaaga actcttcaa cttcaaaaaa
1381 gccaaagagag tcaaaacaga ggatgaaaaa gagaaagacc cagaggagaa gaaagaagtc
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 1561 cccaacccct cagattttat attttctatt tctctgtgta tttatataaa aatttattaa
 1621 atataaatat cccagggac agaaaccaag gcccagagct cagggcagct gtgctgggtg
 1681 agctcttcca ggagccacct tgccacccat tcttcccggt ottaactttg aaccataaag
 1741 ggtgccaggt ctgggtgaaa gggatacttt tatgcaacca taagacaaac tcctgaaatg
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 1861 ctgggtcttc aaggatctcc tgtttttttc aggctcctaa agtaacatca gccattttta
 1921 gattggttct gttttcgtag ctccactg gcctcaagtg agccaagaaa cactgcctgc
 1981 cctctgtctg tcttctcta attctgcagg tggagggtgc tagtctagtt tctttttga
 2041 gatactatct tcatttttgt gagcctcttt gtaataaaat ggtacatttc t

FIGURE 10B Continued